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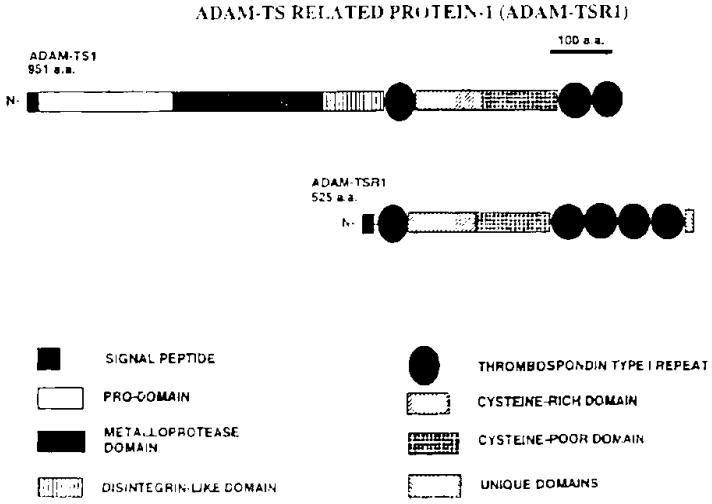
(54) **Title:** NUCLEIC ACIDS ENCODING ZINC METALLOPROTEASES



[illegible] polynucleotides and proteins translated therefrom [illegible] which are human-specific homologs of members of the ADAMTS-N proteins. The present invention also relates to a protein, referred to hereinafter as ADAMTS-R1 (ADAM-TS Related protein-1) and the polynucleotides which encode such protein.

NUCLEIC ACIDS ENCODING ZINC METALLOPROTEASES

Background of the Invention

This invention relates to isolated nucleic acid molecules which encode proteins belonging to a zinc metalloprotease family. The zinc metalloproteases have been implicated in a variety of diseases and development disorders that involve enhanced or depressed proteolysis of components of the extracellular matrix, receptors, or other extracellular molecules.

More particularly, the invention relates to isolated nucleic acid molecules encoding proteins belonging to a subfamily of zinc metalloproteases referred to as "ADAMTS", an abbreviation for A Disintegrin-like And Metalloprotease domain with ThromboSpondin type I motifs. Proteins in the ADAMTS subfamily all possess a Zn protease catalytic site consensus sequence (HEXXH+H), which suggests an intact catalytic activity for each of these proteins. The ADAMTS proteins also have putative N-terminal signal peptides and lack transmembrane domains, which suggests that the proteins in this subfamily are secreted. The proteins in the ADAMTS subfamily also possess at least one thrombospondin type (TSP1) motif, which suggests a binding of these proteins to components of the extracellular matrix (ECM) or to cell surface components.

Members of the ADAMTS subfamily of proteins are ADAMTS-1, ADAMTS-2, ADAMTS-3, and ADAMTS-4. ADAMTS-1 protein is selectively expressed in colon 26 adenocarcinoma cachexigenic sublines. ADAMTS-1 mRNA is induced by the inflammatory cytokine interleukin-1 in vitro and by intravenous administration of lipopolysaccharide in vivo. Thus, the ADAMTS-1 protein is believed to play a role in tumor

cleavage of native triple-helical procollagen I and procollagen II. The ADAMTS-2 protein also has an affinity for collagen XIV. Lack of the ADAMTS-2 protein is known to cause dermatosparaxis in cattle, or Ehlers-Danlos syndrome type VIIC (EDS-VIIC) in humans. EDS-VIIC is characterized clinically by severe skin fragility, and biochemically by the presence in skin of procollagen which is incompletely processed at the amino terminus. Thus, it is believed that the ADAMTS-2 protein plays a role in processing of procollagen to mature collagen, an essential step for correct assembly of collagen into collagen fibrils. The ADAMTS-3 protein is similar in sequence to ADAMTS-2 and may have similar function.

The ADAMTS-4 protein catalyzes cleavage of the core protein of the extracellular matrix proteoglycan, aggrecan. Aggrecan degradation is an important factor in the erosion of articular cartilage in arthritic disease. Aggrecan fragments have been identified in cultures undergoing cartilage matrix degradation and in arthritic synovial fluids. Therefore, overexpression or activation of ADAMTS-4 protein may be related to both inflammatory and non-inflammatory arthritis.

On the basis of the structure, location, and the demonstrated proteolytic activity of ADAMTS proteins 1-4, it is expected that other members of the ADAMTS subfamily play a role in the cleavage of proteoglycan core proteins that are found in the extracellular matrix, such as, for example, versican, brevican, neuracan, NG-2, aggrecan, as well as molecules such as collagen. It is also expected that other members of the ADAMTS subfamily play a role in embryogenesis, implantation of a fertilized egg, angiogenesis

Thus, it is believed that other members of the ADAMTS

subfamily of proteins, the nucleic acids that encode such proteins, and antibodies that are specific for such proteins. Such molecules are useful research tools for studying development of the extracellular matrix during embryogenesis and fetal development, and for studying disorders or diseases that are characterized by improper development of the extracellular matrix or enhanced or reduced destruction of the extracellular matrix. Such molecules, particularly the nucleic acids and the antibodies, are also useful tools for diagnosing such diseases or for monitoring the efficacy of therapeutic agents that have been developed to treat such diseases.

Summary of the Invention

The present invention provides novel, isolated, and substantially purified proteins having the characteristics of an ADAMTS protein. The novel proteins are referred to hereinafter individually as "ADAMTS-5", "ADAMTS-6", "ADAMTS-7", "ADAMTS-8", "ADAMTS-9" and "ADAMTS-10", and collectively as "ADAMTS-N". In one embodiment, the ADAMTS-5 protein is a mature mouse protein which comprises amino acid 231 through amino acid 930 of the sequence set forth set forth in SEQ ID NO: 2. In another embodiment, ADAMTS-5 is a human ADAMTS-5 protein which comprises amino acid 1 through amino acid 518 of the sequence set forth in SEQ ID NO: 4. In one embodiment, mature human ADAMTS-6 protein comprises amino acid 245 through amino acid 860 of SEQ ID NO: 6. In one embodiment, mature human ADAMTS-7 protein comprises amino acid 233 through amino acid 997 of the sequence set forth in SEQ ID NO: 8. In one embodiment, mature ADAMTS-8 protein is a mouse protein which comprises amino acid

[illegible] forth in SEQ ID NO: 10. In one embodiment, mature ADAMTS-9 protein

is a human protein which comprises amino acid 236 through amino acid 1882 of the sequence set forth in SEQ ID NO: 14. In another embodiment, ADAMTS-9 protein is a mouse protein which comprises amino acid 1 through amino acid 974 of the sequence set forth in SEQ ID NO: 16. In one embodiment, mature ADAMTS 10 protein is a human protein which comprises amino acid 212 through amino acid 1081 of the sequence set forth in SEQ ID NO: 18. In another embodiment, ADAMTS-10 protein is a mouse protein which comprises amino acid 1 through amino acid 547 of the sequence set forth in SEQ ID NO: 20

The present invention also provides isolated polynucleotides which encode an ADAMTS-N protein or a variant thereof, polynucleotide sequences complementary to such polynucleotides, vectors containing such polynucleotides, and host cells transformed or transfected with such vectors. The present invention also relates to antibodies which are immunospecific for one or more of the ADAMTS-N proteins. The present invention also relates to a protein referred to hereinafter as ADAMTS-R1 (ADAM-T-S Related protein-1) and the polynucleotides which encode such protein. In one embodiment, the ADAMTS-R1 protein comprises amino acid 1 through amino acid 525 of the sequence set forth in SEQ. ID NO: 22.

Brief Description of the Drawings

Figure 1 shows an amino acid sequence (SEQ ID NO:2) of a full-length mouse ADAMTS-5 protein and a nucleic acid sequence (SEQ ID NO: 1) which encodes such protein.

Figure 2 shows an amino acid sequence (SEQ ID NO:4) of a partial human ADAMTS-5 protein and a nucleic acid sequence (SEQ ID NO: 3) which encodes such protein.

Figure 4 shows an amino acid sequence (SEQ ID NO:8) of a full-length human ADAMTS-7 protein and a nucleic acid sequence (SEQ ID NO:7) which encodes such protein.

Figure 5 shows an amino acid sequence (SEQ ID NO 10) of a full-length mouse ADAMTS-8 protein and a nucleic acid sequence (SEQ ID NO:9) which encodes such protein.

Figure 6 shows an amino acid sequence (SEQ ID NO: 12) of a partial human ADAMTS-8 protein and a nucleic acid sequence (SEQ ID NO: 11) which encodes such amino acid sequence.

Figure 7 shows an amino acid sequence (SEQ ID NO: 14), of a full-length human ADAMTS-9 protein and a nucleic acid sequence (SEQ ID NO: 13) Which encodes such protein.

Figure 8 shows an amino acid sequence (SEQ ID NO. 16) of a partial mouse ADAMTS-9 protein and a nucleic acid sequence (SEQ ID NO: 15) which encodes such amino acid sequence.

Figure 9 shows an amino acid sequence (SEQ ID NO 18) of a full-length human ADAMTS-10 protein and a nucleic acid sequence (SEQ ID NO: 17) which encodes such protein.

Figure 10 shows an amino acid sequence (SEQ ID NO:20) of a partial mouse ADAMTS-10 protein and a nucleic acid sequence (SEQ ID NO: 19) which encodes such amino acid sequence.

Figure 11 shows an amino acid sequence (SEQ ID NO:22) of a full length ADAMTS-B1 protein and a nucleic acid sequence (SEQ ID NO: 21) which encodes such protein.

Figure 12 depicts the cloning strategy used for isolation of **a.** mouse and human ADAMTS-5 cDNAs **b.** human ADAMTS-6 cDNA and **c.** human ADAMTS-7

[illegible] regions of [illegible] alternatively spliced transcripts are shown by the dotted

dotted lines. DNA scale marker (in bp) and amino acid scale marker are at upper right. Location of the probe used for in situ hybridization (ISH) is shown in a.

Figure 13 shows the predicted amino acid sequences of **a.** the mouse and human ADAMTS-5 proteins (alignment shows mouse sequence above, partial human sequence below) **b.** ADAMTS-6, and **c.** ADAMTS-7. The active-site sequences and proposed Met-turn are enclosed in boxes. Potential furin cleavage site(s) are indicated by arrows. Thrombospondin type 1 modules are underlined. Potential sites for N-linked glycosylation are overlined. Cysteine residues within the context of an MMP-like "cysteine switch" are indicated by the solid circles. Other cysteine residues are indicated by asterisks. The prodomain extends until the furin cleavage site, and the catalytic domain extends from the furin cleavage site to the approximate start of the disintegrin-like sequence (Dis). The start of the spacer domain is indicated; the region between the N-terminal TS domain and the spacer domain is the cysteine-rich domain. The single letter amino acid code is used.

Figure 14 shows Northern analysis of expression of ADAMTS-5, 6 and 7. RNA kilobase markers are shown at left of each autoradiogram, and tissue origin is indicated above each lane. **a.** Mouse embryo northern blots. b. Human multiple adult tissue northern blots.

Figure 15 is a schematic representation of the domain structure of ADAMTS-R1 protein as compared to ADAMTS-1 protein.

Figure 16 shows an amino acid sequence (SEQ ID NO: 24) of an alternative embodiment of a full length human ADAMTS-16 protein and a

[illegible] protein designated as human ADAMTS-5a and a nucleic acid sequence

(SEQ ID NO: 25) which encodes such protein.

Figure 18 is a schematic representation of the domain structure of human ADAMTS-9b protein as compared to human and mouse ADAMTS-9 protein.

Detailed Description of the Invention

ADAMTS-N Proteins

The present invention relates to novel, isolated, substantially purified, mammalian proteins belonging to the ADAMTS subfamily of metalloproteases. As used herein, the term "substantially purified" refers to a protein that is removed from its natural environment, isolated or separated, and at least 60% free, preferably 75% free, and most preferably 90% free from other components with which it is naturally associated.

The novel mammalian proteins are ADAMTS-5, ADAMTS-6, ADAMTS-7, ADAMTS-8, ADAMTS-9 and ADAMTS-10, collectively ADAMTS-N. In one embodiment, the ADAMTS-5 protein is a mature mouse protein which comprises amino acid 231 through amino acid 930 of the sequence set forth in SEQ ID NO: 2. In another embodiment, the ADAMTS-5 protein is a human protein which comprises amino acid 1 through amino acid 518 of the sequence set forth in SEQ ID NO: 4. In one embodiment, ADAMTS-6 protein is a mat-Lire human protein which comprises amino acid 245 through amino acid 860 of SEQ ID NO:6. In one embodiment, the ADAMTS-7 protein is a mature human protein which comprises amino acid 233 through amino acid 997 of the sequence set forth in SEQ ID NO: 8. In one embodiment, the ADAMTS-8 protein is a mature mouse protein which comprises amino acid 229 through amino acid 905 of the sequence set forth in SEQ ID NO: 10. In another embodiment, the

[illegible] comprises amino acid [illegible] through amino acid [illegible] of the sequence set

forth in SEQ ID NO: 14. In another embodiment, the ADAMTS-9 protein is a mouse protein which comprises amino acid 1 through amino acid 874 of the sequence set forth in SEQ ID NO: 16. In another embodiment, the ADAMTS-9 designated ADAMTS-9b is a human protein which is comprised of 1934 amino acids as set forth in SEQ ID NO 26. In one embodiment, the ADAMTS-10 protein is a mature human protein which comprises amino acid 212 through amino acid 1081 of the sequence set forth in SEQ ID NO: 18. In another embodiment the ADAMTS- 10 protein is a mouse protein which comprises amino acid 1 through amino acid 525 of the sequence set forth in SEQ ID NO:20. In another embodiment, the ADAMTS-10 protein is a human protein which is comprised of 1072 amino acids as set forth in SEQ ID NO 24.

All of the novel ADAMTS-N proteins starting at the amino terminus comprise a signal sequence followed by a putative pro region which contains a consensus sequence for furin cleavage (except for ADAMTS-10), a catalytic domain, a domain of 60-90 residues with 35 to 45% similarity to snake venom disintegrins, a TS module, a cysteine rich domain containing multiple conserved cysteine residues, a spacer domain, and one or multiple C terminal TS modules. (See Figure 12.) As determined using the BLAST software from the National Center for Biotechnology Information, the predicted mature forms of the ADAMTS-N proteins show an overall 20-30% similarity to each other and to ADAMTS-1-4, although this may be considerably higher or lower for individual domains as described below.

The ADAMTS-N proteins also encompass variants of the ADAMTS-N proteins shown in Figs. 1-10. A "variant" as used herein, refers to

of the reference sequence. The variant protein has an altered sequence

in which one or more of the amino acids in the reference sequence is deleted or substituted, or one or more amino acids are inserted into the sequence of the reference amino acid sequence. As a result of the alterations, the variant protein has an amino acid sequence which is at least 95% identical to the reference sequence, preferably, at least 97% identical, more preferably at least 98% identical, most preferably at least 99% identical to the reference sequence. Variant sequences which are at least 95% identical have no more than 5 alterations, i.e. any combination of deletions, insertions or substitutions, per 100 amino acids of the reference sequence. Percent identity is determined by comparing the amino acid sequence of the variant with the reference sequence using MEGALIGN project in the DNA STAR program. Sequences are aligned for identity calculations using the method of the software basic local alignment search tool in the BLAST network service (the National Center for Biotechnology Information, Bethesda, MD) which employs the method of Altschul, S. F., Gish, W., Miller, W., Myers, E. W. & Lipman, D. J. (1990) *J. Mol. Biol.* 215, 403-410. Identities are calculated by the Align program (DNAstar, Inc.) In all cases, internal gaps and amino acid insertions in the candidate sequence as aligned are not ignored when making the identity calculation.

While it is possible to have nonconservative amino acid substitutions, it is preferred that the substitutions be conservative amino acid substitutions, in which the substituted amino acid has similar structural or chemical properties with the corresponding amino acid in the reference sequence. By way of example,

. . . . .

. amino acid e.g. serine and threonine with another substitution at

one acidic residue, e.g. glutamic acid or aspartic acid, with another; replacement of one amide-containing residue, e.g. asparagine and glutamine, with another; replacement of one aromatic, residue, e.g. phenylalanine and tyrosine, with another; replacement of one basic residue, e.g. lysine, arginine and histidine, with another; and replacement of one small amino acid, e.g., alanine, serine, threonine, methionine, and glycine, with another.

The alterations are designed not to abolish the immunoreactivity of the variant protein with antibodies that bind to the reference protein. Guidance in determining which amino acid residues may be substituted, inserted or deleted without abolishing immunoreactivity of the variant protein with an antibody specific for the respective reference protein are found using computer programs well known in the art, for example, DNASTAR software.

The ADAMTS-N proteins also encompass fusion proteins comprising an ADAMTS-N protein and a tag, i.e., a second protein or one or more amino acids, preferably from about 2 to 65 amino acids, more preferably from about 34 to about 62 amino acids, which are added to the amino terminus of, the carboxy terminus of, or any point within the amino acid sequence of an ADAMTS-N protein, or a variant of such protein. Typically, such additions are made to stabilize the resulting fusion protein or to simplify purification of an expressed recombinant form of the corresponding ADAMTS-N protein or variant of such protein. Such tags are known in the art. Representative examples of such tags include sequences which encode a series of histidine residues, the epitope tag FLAG, the Herpes simplex

[illegible] amino acids, preferably no more than 1 amino acid at

the respective ADAMTS-N protein are altered by posttranslation processes or synthetic methods. Examples of such modifications include, but are not limited to, acetylation, amidation, ADP-ribosylation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or a lipid, cross-linking gamma-carboxylation, glycosylation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, sulfation, and transfer-RNA mediated additions of amino acids to proteins such as arginylation and ubiquitination.

The ADAMTS-N proteins are immunogenic and, thus, are useful for preparing antibodies. Such antibodies are useful for identifying and diagnosing disorders which are associated with decreased expression or activity or increased expression of an ADAMTS-N protein. The ADAMTS-N protein may also be useful for treating such disorder.

Diseases involving enhanced or depressed proteolyisis of the core proteins of the extracellular may involve enhanced expression or activity or decreased expression or activity of one or more ADAMTS-N proteins. Thus, ADAMTS-N proteins may be used to identify drugs, polypeptides, auto-antibodies, or other natural compounds which bind to an ADAMTS-N protein with sufficient affinity to block or facilitate its activity. The activity of the ADAMTS-N protein is assayed in the presence and the absence of the putative inhibitor or facilitator using any of a variety of protease assays known in the art. In general, the activity of the ADAMTS-N protein is assayed through the use of a peptide or protein substrate having a known or

example, the substrate may be tagged with a fluorescent group on one

side of the cleavage site and with a fluorescence-quenching group on the opposite side of the cleavage site. Upon cleavage by the substrate, quenching is eliminated and a detectable signal is produced. Alternatively, the substrate is tagged with a colorimetric leaving group that more strongly absorbs upon cleavage. Agents which block ADAMTS-N-catalyzed cleavage of a protein substrate may be administered to a subject to block proteolysis of the corresponding protein substrate.

ADAMTS-R1 Protein

The present invention also relates to a protein, referred to hereinafter as "ADAMTS-R1". From its amino to its carboxyl terminus, ADAMTS-R1 comprises a signal peptide sequence, a TS1 module, a cysteine-rich domain, a spacer domain, and three TS1 modules. Thus, ADAMTS-R1 has a structure which is related to or similar to an ADAMTS-N protein, but which lacks a catalytic domain and a disintegrin-like domain. In one embodiment, ADAMTS-R1, protein comprises amino acid 1 through amino acid 525 of the amino acid sequence, SEQ ID NO:22, shown in Fig. 11. Such protein has a 30-40% overall sequence identity with similar regions of the ADAMTS-N proteins. The ADAMTS-R1 proteins also encompass variants of the amino acid sequence shown in Fig. 11 and fusion proteins which contain the amino acid sequence shown in Fig. 11 or a variant thereof. On the basis of its domain organization, it is expected that ADAMTS-R1 can bind to extracellular matrix or cell surface molecules, including ADAMTS-N substrates. Thus, it is expected that ADAMTS-R1 can be used as an cell-matrix or cell-cell adhesion molecule or an ADAMTS-N competitive inhibitor. The ADAMTS-R1

expression or increased expression of an ADAMTS-R1 protein.

Polynucleotides

The present invention also provides isolated polynucleotides which encode the mammalian ADAMTS-N proteins and the mammalian ADAMTS-R1 protein. Figure 1 shows one embodiment of a polynucleotide, SEQ ID NO: 1, which encodes the full-length mouse ADAMTS-5 protein. Figure 2 shows one embodiment of a polynucleotide; SEQ ID NO: 3, which encodes a partial human ADAMTS-5 protein. Figure 3 shows one embodiment of a polynucleotide; SEQ ID NO: 5, which encodes a full-length human ADAMTS-6 protein. Figure 4 shows one embodiment of a polynucleotide; SEQ ID NO: 7, which encodes a full-length human ADAMTS-7 protein. Figure 5 shows one embodiment of a polynucleotide; SEQ ID NO: 9, which encodes a full-length mouse ADAMTS-8 protein. Figure 6 shows one embodiment of a polynucleotide; SEQ ID NO: 11, which encodes a partial human ADAMTS-8 protein. Figure 7 shows one embodiment of a polynucleotide; SEQ ID NO: 13, which encodes a full-length human ADAMTS-9 protein. Figure 8 shows one embodiment of a polynucleotide; SEQ ID NO: 15, which encodes a partial ADAMTS-9 protein. Figure 9 shows one embodiment of a polynucleotide; SEQ ID NO: 17, which encodes a full-length human ADAMTS-10 protein. Figure 10 shows one embodiment of a polynucleotide; SEQ ID NO: 19, which encodes a partial mouse ADAMTS-10 protein. Figure 11 shows one embodiment of a polynucleotide; SEQ ID NO: 21, which encodes a full-length ADAMTS-R1 protein.

Due to the known degeneracy of the genetic code wherein more than one codon can encode the same amino acid, a DNA sequence may vary from that shown in SEQ ID NO: 1 and still encode an ADAMTS-5

in SEQ ID NO:6. Similarly a DNA sequence may vary from that shown in SEQ ID NOS: 7, 9, 11, and 13, and still encode the amino acid sequences shown in SEQ ID NOS: 8, 10, 12, and 14, respectively. Such variant DNA sequence may result from silent mutations, such as for example those that occur during PCR amplification or from deliberate mutagenesis of a native sequence.

The present polynucleotides also encompass polynucleotides having sequences that are capable of hybridizing to the nucleotide sequences of FIGS 1 - 11 under stringent conditions, preferably highly stringent conditions. Hybridization conditions are based on the melting temperature™ of the nucleic acid binding complex or probe, as described in Berger and Kimmel (1987) Guide to Molecular Cloning Techniques, Methods in Enzymology, vol 152, Academic Press. The term "stringent conditions, as used herein, is the "stringency" which occurs within a range from about Tm-5 (5° below the melting temperature of the probe) to about 20° C below Tm. As used herein "highly stringent" conditions employ at least 0.2 x SSC buffer and at least 65° C. As recognized in the art, stringency conditions can be attained by varying a number of factors such as the length and nature, i.e., DNA or RNA, of the probe; the length and nature of the target sequence, the concentration of the salts and other components, such as formamide, dextran sulfate, and polyethylene glycol, of the hybridization solution. All of these factors may be varied to generate conditions of stringency which are equivalent to the conditions listed above.

The present polynucleotides also encompasses alleles of the

[illegible] may result from one or more mutations in the [illegible]

ADAMTS-R1 encoding sequence. Such mutations typically arise from natural addition, deletion of substitution of nucleotides in the open reading frame sequences. Any gene which encodes an ADAMTS-N protein or ADAMTS-RI protein may have none, one, or several allelic forms. Such alleles are identified using conventional techniques, such as for example screening libraries with probes having sequences identical to or complementary with one or more ADAMTS-N polynucleotides.

The present polynucleotides also encompass altered polynucleotides which encode ADAMTS-N proteins, ADAMTS-R1 proteins, and variants thereof. Such alterations include deletions, additions, or substitutions. Such alterations may produce a silent change and result in an ADAMTS-N protein having the same amino acid sequence as the ADAMTS-N protein encoded by the unaltered polynucleotide. Such alterations may produce a nucleotide sequence possessing non-naturally occurring codons. For example, codons preferred by a particular prokaryotic or eucaryotic host may be incorporated into the nucleotide sequences showing Figures 1 -11 to increase the rate of expression of the proteins encoded by such sequences. Such alterations may also introduce new restriction sites into the sequence or result in the production of an ADAMTS-N or ADAMTS-PI variant. Typically, such alterations are accomplished using site-directed mutagenesis.

The polynucleotides are useful for producing ADAMTS-N or ADAMTS-R1 proteins. For example, an RNA molecule encoding an ADAMTS-N protein is used in a cell-free translation systems to prepare such

[illegible] recombinant and synthetic DNA [illegible]

SV40, bacterial plasmids, phage DNAs; yeast plasmids, vectors derived from combinations of plasmids and phage DNAs, viral DNA such as vaccinia, adenovirus, fowl pox virus, pseudorabies, baculovirus, and retrovirus. The DNA sequence is introduced into the expression vector by 5 conventional procedures.

Accordingly, the present invention also relates to recombinant constructs comprising one or more of the present polynucleotide sequences. Suitable constructs include, for example, vectors, such as a plasmid, phagemid, or viral vector, into which a sequence that encodes an ADAMTS-N protein or an ADAMTS-R1 protein has been inserted. In the expression vector, the DNA sequence which encodes the ADAMTS-N protein is operatively linked to an expression control sequence, i.e., a promoter, which directs mRNA synthesis. Representative examples of such promoters, include the LTR or SV40 promoter, the *E. coli* lac or trp, the phage lambda PL promoter and other promoters known to control expression of genes in prokaryotic or eukaryotic cells or in viruses. The promoter may also be the natural promoter of the ADAMTS-N encoding sequence. The expression vector, preferably, also contains a ribosome binding site for translation initiation and a transcription terminator. Preferably, the recombinant expression vectors also include an origin of replication and a selectable marker, such as for example, the ampicillin resistance gene of *E. coli* to permit selection of transformed cells, i.e. cells that are expressing the heterologous DNA sequences. The polynucleotide sequence encoding the ADAMTS-N protein is incorporated into the vector in frame with translation

[illegible] art. Such techniques are described in Sambrook, J. et al

(1989) Molecular Cloning A Laboratory Manual, Cold Spring Harbor Press, Plainview, N.Y. and Ausubel, F. M. et al. (1989) Current Protocols in Molecular Biology, John Wile & Sons, New York, NY.

Polynucleotides encoding an ADAMTS-N or ADAMTS-R1 protein may also be used for diagnostic purposes. The polynucleotides may be used to detect and quantify ADAMTS-N or ADAMTS-R1 gene transcripts in biopsied tissues in which enhanced expression or reduced expression of the corresponding ADAMTS-N or ADAMTS-RI gene is correlated with a disease. The diagnostic assay may be used to determine whether expression is absent, present, or altered and to determine whether certain therapeutic agents modulate expression of the corresponding ADAMTS-N or ADAMTS-R1 gene.

Also encompassed by the present invention, are single stranded polynucleotides, hereinafter referred to as antisense polynucleotides, having sequences which are complementary to the DNA and RNA sequences which encode the ADAMTS-N or ADAMTS-R1 proteins. The term complementary as used herein refers to the natural binding of the polynucleotides under permissive salt and 5 temperature conditions by base pairing.

The present invention also encompasses oligonucleotides that are used as primers in polymerase chain reaction (PCR technologies to amplify transcripts of the genes which encode the ADAMTS-N and ADAMTSR-1 proteins or portions of such transcripts. Preferably, the primers comprise 18-30 nucleotides, more preferably 19-25 nucleotides. Preferably, the primers have a G-C content of 40% or greater. Such oligonucleotides are at least 98% complementary with a

[illegible] complementary, more preferably [illegible] complementary, with a

sense strand or its corresponding antisense strand. Primers which are which have 100% complementarity with the antisense strand of a double-stranded DNA molecule which encodes an ADAMTS-N protein have a sequence which is identical to a sequence contained within the sense strand. The identity of primers which are 15 nucleotides in length and have full complementarity with a portion of the antisense strand of a double-stranded DNA molecule which encodes the ADAMTS-N protein is determined using the nucleotide sequences, shown in FIG I - 11 and described by the general formula a-b; where a is any integer between I and the position number of the nucleotide which is located 15 residues upstream of the 3' end of the sense or antisense strand of the cDNA sequences shown in FIG 1 -11; where b is equal to a+14; and where both a and b correspond to the positions of nucleotide residues of the cDNA sequences shown in FIGS 1 - 11.

The present invention also encompasses oligonucleotides that are useful as hybridization probes for for isolating and identifying cDNA clones and genomic clones encoding the ADAMTS-N or ADAMTS-R1 protein or allelic forms thereof. Such hybridization probes are also useful for detecting transcripts of the genes which encode the ADAMTS-N family proteins or for mapping of the genes which encode the ADAMTS-N proteins Preferably, such oligonucleotides comprise at least 210 nucleotides, more preferably at least 230, most preferably from about 210 to 280 nucleotides. Such hybridization probes have a sequence which is at least 90% complementary with a sequence contained within the sense strand of a DNA molecule which encodes an ADAMTS-N protein or ADAMTS-R1 protein or with a sequence contained

within a range from about [illegible] with the [illegible]

Tm of the probe) to about 20°C to 25°C below Tm. The probes are used in Northern assays to detect transcripts of ADAMTS-N homologous genes and in Southern assays to detect ADAMTS-N homologous genes. The identity of probes which are 200 nucleotides 5 in length and have full complementarity with a portion of the antisense strand of a double-stranded DNA molecule which encodes the ADAMTS-N protein is determined using the nucleotide sequences shown in FIG 1 - 10 and described by the general formula a-b; where a is any integer between I and the position number of the nucleotide which is located 200 residues upstream of the 3' end of the sense or antisense strand of the cDNA sequences shown in FIG 1 -10; b is equal to a +200; and where both a and b correspond to the positions of nucleotide residues of the cDNA sequences shown in FIG 1-10.

Such probes or primers are also useful for identifying tissues or cells in which the corresponding ADAMTS-N or ADAMTS-R1 gene is preferentially expressed either constitutively or at particular state of tissue differentiation or development or in disease states. Expression of the ADAMTS-N or ADAMTS-R1 gene in a particular tissue or group of cells is determined using conventional procedures including, but not limited to, Northern analysis, in situ hybridization to RNA or RT-PCR amplification. Isolated polynucleotides encoding an ADAMTS-N or ADAMTS-R1 protein are also useful as chromosome markers to map linked gene positions to identify chromosomal aberrations such as translocations, inversions and trisomies, to compare with endogenous DNA sequences in patients to identify potential genetic disorders, and as probes to hybridize

and chromosomes, PCR, and allele specific hybridization.

Antibodies

In another aspect, the present invention relates to antibodies which are specific for and bind to the ADAMTS-5 protein, the ADAMTS-6 protein, the ADAMTS-7 protein, the ADAMTS-8 protein, the ADAMTS-9 protein, the ADAMTS-10 protein, or the ADAMTS-R1 protein. Such antibodies are useful research tools for identifying *tissues that contain elevated levels of the respective protein and for purifying the respective protein from cell or tissue extracts, medium of cultured cells, or partially purified preparations of intracellular and extracellular proteins by affinity chromatography. Such antibodies are also useful for identifying and diagnosing diseases associated with elevated or reduced levels of an ADAMTS-N protein or ADAMTS-R1 protein. Such antibodies are also useful for monitoring the effect of therapeutic agents on the synthesis and secretion of ADAMTS-N proteins by cells in vitro and in vivo. Such antibodies may also be employed in procedures, such as co-immunoprecipitation and co-affinity chromatography, for identifying other proteins, activators and inhibitors which bind to an ADAMTS-N or ADAMTS-R1 protein.

The present invention also provides a method for detecting an ADAMTS-N or ADAMTS-R1 protein, in a bodily sample from a patient using antibodies immunospecific for an ADAMTS-N or ADAMTS-R1 protein. The method comprises contacting the antibody with a sample taken from the patient; and assaying for the formation of a complex between the antibody and the corresponding ADAMTS-N or ADAMTS-R1 protein present in the sample. The sample may be a tissue or a biological fluid,

tissue, cells obtained from swabs and smears. To monitor changes in expression of the ADAMTS-N protein during fetal development and pregnancy, it is preferred that the sample be amniotic fluid. To monitor changes in expression of the ADAMTS-N protein during joint disorders, the preferred sample is synovial fluid. To monitor changes in expression of ADAMTS-N proteins during cancer, the preferred samples include, but are not limited to, serum, body fluids, or biopsy tissue. To monitor changes in expression of ADAMTS-N proteins during inflammation the preferred samples include, but are not limited to, serum, body fluids, or biopsy tissue.

The sample may be untreated, or subjected to precipitation; fractionation, separation, or purification before combining with the anti-ADAMTS-N protein antibody. For ease of detection, it is preferred that isolated proteins from the sample be attached to a substrate such as. a column, plastic dish, matrix, or membrane, preferably nitrocellulose. Preferably, the detection method employs an enzyme-linked immunosorbent assay (ELISA) or a Western immunoblot procedure.

Interactions between an ADAMTS-N protein in the sample and the corresponding anti ADAMTS-N antibody are detected by radiometric, colorimetric or fluorometric means, size separation, or precipitation. Preferably, detection of the antibody-ADAMTS-N protein complex is by addition of a secondary antibody that is coupled to a detectable tag, such as for example, an enzyme, fluorophore, or chromophore. Formation of the complex is indicative of the presence of the ADAMTS-N protein i n the test sample. Thus,

[illegible] to quantify the amount of the ADAMTS-N protein in the test sample.

Deviation between control and test values establishes the parameters for diagnosing the disease.

Preparing the ADAMTS-N proteins and the ADAMTS-R1 protein

The ADAMTS-N proteins and the ADAMT-SR1 protein may be produced by conventional peptide synthesizers. The ADAMTS-N proteins and the ADAMTS-R1 protein may also be produced using cell-free translationsystems and RNA molecules derived from DNA constructs that encode an ADAMTS-N protein or an ADAMTS-RI protein. Alternatively, ADAMTS-N proteins are made by transfecting host cells with expression vectors that comprise a DNA sequence that encodes the respective ADAMTS-N protein and then inducing expression of the protein in the host. cells. For recombinant production, recombinant constructs comprising one or more of the sequences which encode the ADAMTS-N protein or a variant thereof are introduced into host cells by conventional methods such as calcium phosphate transfection, DEAE-dextran mediated transfection, transvection, microinjection, cationic lipid-mediated transfection, electroporation, transduction, scrape lading, ballistic introduction or infection.

The ADAMTS-N protein and the ADAMTS-R1 protein may be expressed in suitable host cells, such as for example, mammalian cells, yeast, bacteria, insect cells or other cells under the control of appropriate promoters using conventional techniques. Suitable hosts include, but are not limited to, E. coli, P. pastoris, Cos cells and 293 HEK cells. Following transformation of the suitable host strain and growth of the host strain to an appropriate cell density, the cells are harvested by centrifugation, disrupted by physical or

[illegible] transformed host cells, such as [illegible]

cell pellets or from cell culture medium, followed by salting-out, and one or more chromatography steps, including aqueous ion exchange chromatography, size exclusion chromatography steps, and high performance liquid chromatography (HPLC), and affinity chromatography may be used to isolate the recombinant ADAMTS-N protein or ADAMTS R1 protein

Preparation of Antibodies

The ADAMTS-N proteins, and variants thereof are used as immunogens to produce antibodies immunospecific for one or more ADAMTS-N protein. The term "immunospecific" means the antibodies have substantially greater affinity for one or more ADAMTS-N protein than for other proteins. Such antibodies may include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, and Fab fragments.

Antibodies are also prepared using an oligopeptide having a sequence which is identical to a portion of the amino acid sequence of an ADAMTS-N protein. Preferably the oligopeptide has an amino acid sequence of at least five amino acids, and more preferably, at least 10 amino acids that are identical to a portion of the amino acid sequence of an ADAMTS-N protein. Such peptides are conventionally fused with those of another protein such as keyhole limpet hemocyanin and antibody produced against the chimeric molecule. One preferred oligopeptide for preparing an antibody to mouse ADAMTS-5 has the sequence (C)HIRVRQFKAHQTRF, SEQ ID NO: 30. Another preferred oligopeptide for preparing an antibody to ADAMTS-5 is CEAKNGYQSDAKGVKTFVEWVPKYAG, SEQ ID NO: 3 1. One preferred oligopeptide for preparing an antibody to ADAMTS-6 has the sequence

preparing an antibody to ADAMTS-8 has the sequence CVKEDVENPKAVVDGDWGP, SEQ ID NO:25. One preferred oligopeptide for preparing an antibody to ADAMTS-9 has the sequence QHPFQNEDYRPRSASPSRTH, SEQ ID NO:26. Another preferred oligopeptide for preparing an antibody to ADAMTS-9 has the sequence PQNCKEVKRLKGASEDGEYF, SEQ ID NO:27. One preferred oligopeptide for preparing an antibody for ADAMTS-R1 has the sequence QELEEGAAVSEEPS, SEQ ID NO:28. Another preferred oligopeptide for preparing an antibody for ADAMTS-R1 has the sequence YYPENIKPKPKLQE; SEQ ID NO:29.

Polyclonal antibodies are generated using conventional techniques by administering the ADAMTS-N protein or achimeric molecule to a host animal. Depending on the host species, various adjuvants may be used to increase immunological response. Among adjuvants used in humans, Bacilli Calmette-Guerin (BCG), and Corynebacterium parvum. are especially preferable. Conventional protocols are also used to collect blood from the immunized animals and to isolate the serum and or the IgG fraction from the blood.

For preparation of monoclonal antibodies, conventional hybridoma techniques are used. Such antibodies are produced by continuous cell lines in culture. Suitable techniques for preparing monoclonal antibodies include, but are not limited to, the hybridoma technique, the human B-cell hybridoma technique, and the EBV hybridoma technique.

Various immunoassays may be used for screening to identify antibodies having the desired specificity. These include protocols which involve competitive binding or immunoradiometric assays and

Polynucleotides encoding antibodies to ADAMTS-N

protein or an ADAMTS-R1 protein may be synthesized in whole or in part using chemical methods. Polynucleotides which encode an ADAMTS-N protein, particularly alleles of the genes which encode the ADAMTS-N protein, may be obtained by screening a genomic library or cDNA library with a probe comprising sequences identical or complementary to the sequences shown in Figures 1 - 10 or with antibodies immunospecific for a ADAMTS-N protein to identify clones containing such polynucleotide.

Example 1 ADAMTS-512 protein

A cDNA encoding mouse ADAMTS-5 protein was obtained using IMAGE Clone 569515, purchased from Research Genetics, Huntsville, Alabama and 7 day old mouse embryo cDNA library from Clontech, Palo Alto, CA. A cDNA encoding human ADAMTS-5 protein was obtained using IMAGE Clone 345484 purchased from Research Genetics, Huntsville, Alabama and a human fetal brain cDNA from Clontech. The clone inserts were sequenced in their entirety. Using oligonucleotide primers based on the sequences at the ends of the clone inserts as template, successive rounds of RACE (Rapid Amplification of cDNA Ends) by PCR was performed at 5' and 3 ends. RACE primers were generated 50-200 bp from the ends of the sequences so that the contiguity of RACE clones with the I.M.A.G.E. clone could be clearly established. A single round of 5' and 3' 20 RACE sufficed for cloning of the entire coding sequence of the mouse ADAMTS-5 protein and part of the catalytic zinc binding site through to the stop codon of the human ADAMTS-5 protein. Primers were designed with calculated Tm>72°C and RACE was performed with nested primers for each amplification. PCR used the Advantage PCR reagents (Clontech, Palo Alto, CA); the

[illegible] RACE used the [illegible]

conditions; 95°C for 1 minute followed by 5 cycles of 95°C for 0.5 minutes, 72°C for 5 minutes, then 5 cycles of 95°C for 0.5 minutes, 70°C for 5 minutes and 20 cycles of 95°C for 0.5 minutes, 68°C for 5 minutes. The PCR products were analyzed by Southern blotting, initially using [α^{32}P]-dCTP labeled.

Hybridizing bands were ligated into pGEM-T Easy (Promega, Madison, WI) and individual clones were selected by another round of Southern analysis. Automated nucleotide sequencing of both strands of each clone were done at the Molecular Biotechnology Core of the Lerner Research Institute, Cleveland Clinic Foundation and nucleotide sequence data were analyzed using the DNAStar software. By integration of the overlapping sequences thus obtained, a contiguous nucleotide sequence was determined. The nucleotide sequence of the mouse ADAMTS-5 cDNA and the predicted amino acid sequence of the protein encoded by this cDNA are shown in Fig. 1. The nucleotide sequence of the human ADAMTS-5 cDNA and the predicted partial amino acid sequence of the protein encoded by this cDNA are shown in Fig. 2.

The predicted molecular mass (Mr) of the mature ADAMTS-5 protein is 73717.50 daltons. It is expected that the actual Mr of the active ADAMTS-5 protein is different due to post-translational modification, which could potentially increase the Mr. The predicted domain organization of ADAMTS-5 protein relative to the cloned cDNA is shown in Figure 12. The pro-domain of the full-length mouse ADAMTS-5 protein has 3 consensus cleavage signals for furin. The most carboxyl-terminal furin cleavage site in ADAMTS-5 predicts the

1. five cysteine residues are upstream of the zinc binding sequence

while three residues are downstream, an arrangement that is shared with other ADAMTS members. The zinc binding signature is followed by a "Met-turn". The catalytic domain is followed by a domain with 35% similarity to snake venom disintegrins. The disintegrin domain contains eight cysteine residues. The first TS repeat contains 52 residues and is followed by a conserved cysteine-rich sequence termed the cysteine-rich domain, designated "CRD", to distinguish it from the cysteine-free spacer domain. The CRD contains ten conserved cysteines and demonstrates high sequence homology with the CRD of other ADAMTS-N proteins. The spacer domain of mouse ADAMTS-5 is 158 amino acids in length and is followed by a second TS module. ADAMTS-5 contains three potential glycosylation sites in the mature protease one of which is just upstream of the start of the spacer domain and the second lies within the spacer domain and the third is near the start of the disintegrin domain. The human ADAMTS-5 protein and the mouse ADAMTS-5 protein have 96% sequence identity. ADAMTS-5 bears 46% sequence identity to ADAMTS-4 (KIAA0688), which is characterized as being involved in catabolism of aggrecan core protein in arthritis and 60% identity to ADAMTS-1 which is involved in inflammation.

Example 2 ADAMTS-6

The nucleotide sequence of a human cDNA encoding the full-length ADAMTS-6 protein was obtained using IMAGE clone 742630, which encodes EST AA400393, and a human fetal brain cDNA from Clontech. RACE was performed as described above in Example 1. The I.M.A.G.E. clone 742630 contained an ORF flanked by consensus splice sequences, indicating the presence of introns. Two successive rounds of RACE at the 5' end and a single round of RACE at the 3' end provided the

The nucleotide sequence of the ADAMTS-6 DNA is shown in Fig. 3 The predicted amino acid sequence, SEQ ID NO:6, of the ADAMTS-6 protein is also shown in Fig. 3. The predicted Mr of the full-length, unprocessed ADAMTS-6 protein is 97,115 daltons., and the predicted Mr of the mature ADAMTS-6 protein is 68412.10 daltons. The domain organization of the ADAMTS-6 protein is shown in Fig. 12. The pro-domain of the full-length ADAMTS-6 protein has one consensus cleavage signal for furin. The catalytic domain of the ADAMTS-6 contains six cysteine residues and the reprolysin -zinc binding signature sequence, HEIVHNFGMNHD, which is followed by a "Met-turn". The catalytic domain is followed by a domain with 35% similarity to snake venom disintegrins. The disintegrin domain contains eight cysteine residues. The first TS repeat contains 52 residues and is followed by a conserve CRD sequence which contains ten conserved cysteines and demonstrates high sequence homology with the CRD of other ADAMTS proteins. The spacer domain of ADAMTS-6 is 127 amino acids in length and is followed by a second TS module. ADAMTS-6 contains four potential glycosylation sites within the pyo-domain and two in the mature protease one of which is in the cysteine rich domain and the other of which is in the spacer domain. ADAMTS-6 bears 46% sequence identity to ADAMTS-1, which is involved in inflammation.

Example 3 ADAMTS-7.

The nucleotide sequence of a cDNA encoding an ADAMTS-7 protein was obtained using IMAGE clone 272098, which encodes EST N4.8032, and a human fetal brain cDNA from Clontech. RACE was performed as described above in Example 1. The I.M.A.G.E. clone 272098 encoded a

methionine codon lies within a satisfactory Kozak consensus for translation initiation.

The nucleotide sequence of the ADAMTS-7 cDNA is shown in Fig. 4. The predicted amino acid sequence, SEQ ID NO: 8, of the ADAMTS-7 protein is also shown in Fig. 4. The predicted Mr of the hill-length, unprocessed ADAMTS-7 protein is 116,607 daltons, and the predicted Mr of the mature ADAMTS-7 protein is 84005 daltons. The domain organization of the ADAMTS-7 protein is shown in Fig. 12. The pro-domain of the full length ADAMTS 7 protein has one consensus cleavage signal for furin. The catalytic domain of the ADAMTS-7 protein contains eight cysteine residues and the reprolysin-zinc binding signature sequence, HELGHSFGIQHD, which is followed by a "Met-turn". The catalytic domain is followed by a domain with 30% similarity to snake venom disintegrins The disintegrin domain contains eight cysteine residues. The first TS repeat contains 52 residues and is followed by a conserved CRD sequence which contains ten conserved cysteines. The spacer domain of ADAMTS-7 is 221 amino acids in length and is followed by a second TS module and a short sequence containing two cysteine residues. ADAMTS-7 contains three potential glycosylation sites within the mature protease; one of which is just upstream of the spacer domain and one of which is within the spacer domain. ADAMTS-7 bears 35 % sequence identity to ADAMTS-1, which is characterized as being involved in inflammation and 32% identity to ADAMTS-2 which is a procollagen processing enzyme.

Example 4: ADAMTS-8

A nucleotide sequence of a cDNA encoding a partial ADAMTS-8 human

protein was obtained using IMAGE clone 2119838, which encodes EST AI400905, and a human fetal brain cDNA library from Clontech. RACE was performed, as described above in Example 1. The nucleotide sequence of the cDNA encoding the full-length ADAMTS-8 mouse protein and the amino acid sequence of such protein is shown in Fig. 5. The nucleotide sequence of the cDNA encoding the partial ADAMTS-8 human protein and the amino acid sequence of such protein is shown in Fig. 6.

The predicted Mr of the full-length, unprocessed ADAMTS-8 mouse protein is 1260693 daltons, and the predicted Mr of the mature ADAMTS-8 protein is 68412.10 daltons. The pro domain of the full-length ADAMTS-8 protein has one consensus cleavage signal for furin. The catalytic domain contains eight cysteine residues and the reprolysm-zinc binding signature sequence, HELGHVLSMPHD, which is followed by a "Met-turn". The catalytic domain is followed by a domain with 20-30% similarity to snake venom disintegrins. The disintegrin-like domain contains eight cysteine residues. The first TS repeat is followed by a conserved CRD sequence which contains 10 conserved cysteines. The spacer domain of ADAMTS-8 is 146 amino acids in length and is followed by a second TS module. The ADAMTS-8 protein contains 4 potential glycosylation sites within the mature protease: one is in the cysteine-rich domain; one is in the catalytic domain; and two are in the disintegrin-like domain. ADAMTS-8 bears 46% sequence identity to ADAMTS-1 and 42% identity to ADAMTS-4.

Example 5: ADAMTS-9

The nucleotide sequence of a cDNA encoding a full-length, human

[illegible] of a cDNA encoding a partial ADAMTS-[illegible]

protein was obtained using IMAGE clone 535663, which encodes EST AA1 06215, and a mouse cDNA library obtained from Clonetech. RACE was performed as described above in Example 1. The nucleotide sequence of the cDNA encoding the full-length ADAMTS-9 human proteinand the amino acid sequence of such protein is shown in Fig.6. The nucleotide sequence of the cDNA encoding the partial ADAMTS-9 mouse protein and the amino acid sequence of such protein is shown in Fig. 7.

The predicted Mr of the mature human ADAMTS-9 protein is 169777.20 daltons. The prodomain of the predicted ADAMTS-9 protein has 3 consensus cleavage signal for furin. The catalytic domain of the ADAMTS-9 contains eight cysteine residues and the reprolysin - zinc binding signature sequence, HELGHVFNMPHD, which is followed by a "Met-turn". The catalytic domain is followed by a domain with 25-30% similarity to snake venom disintegrins The disintegrin domain contains eight cysteine residues. The first TS repeat contains is followed by a conserved CRD sequence which. contains 10 conserved cysteines. The spacer domain of ADAMTS-9 is 124 amino acids in length and is followed by 14 additional TS modules and a C-terminal domain. The ADAMTS-9 protein contains 6 potential glycosylation sites within the mature protease: one in the spacer domain, one in TSP 1 -7, one in TSPI-8, and 3 in the C-terminal domain. The ADAMTS-9 bears 44% sequence identity to ADAMTS-4.

Example 6: ADAMTS-10

The nucleotide sequence of a cDNA encoding a fall-length ADAMTS- 10 protein was obtained using IMAGE clone 110403, which encodes EST AA588434, and a human fetal brain cDNA from Clonetech. The nucleotide sequence of a cDNA encoding a partial, mouse ADAMTS-10 protein was obtained using IMAGE clone 1077653, which encodes EST

performed as described above in Example 1. The nucleotide sequence of the human ADAMTS-10 cDNA and the predicted amino acid sequence, SEQ ID 18, of the human ADAMTS-10 protein encoded by such DNA is shown in Fig. 9. The nucleotide sequence of the cDNA encoding the partial mouse ADAMTS-10 protein and the amino acid sequence of such protein is shown in Fig. 10.

The predicted Mr of the mature ADAMTS-10 protein is 95238 daltons. The pro-domain of the full-length ADAMTS-10 protein has no consensus cleavage signal for furin. The catalytic domain of the ADAMTS-10 contains eight cysteine residues and the reprolysin-zinc binding signature sequence, HEIGHTFGMNHD, which is followed by a "Met-turn". The catalytic domain is followed by a domain with 30% similarity to snake venom disintegrins. The disintegrin-like domain contains eight cysteine residues. The first TS repeat is followed by a conserved CRD sequence which contains 8 conserved cysteines. The spacer domain of ADAMTS-10 is followed by 4 additional TS modules and a Kunitz domain. The ADAMTS-10 protein contains 2 potential glycosylation sites within the mature protease: one in the catalytic domain, and one in the TS 1-3 domain. ADAMTS-10 bears approximately 40% sequence identity to ADAM-TS1, which is characterized as being involved in inflammation.

Comparison of the ADAMTS-N Proteins.

As shown in Figure 11, the ADAMTS-5, ADAMTS-6, and ADAMTS-7 proteins share a common domain organization. From amino to carboxyl termini, they are as follows:

1. **A pre-pro region.** A typical signal sequence of variable length is followed by a putative pro-region of variable length but

context similar to the cysteine "switch" of the MMPs. All three novel cDNAs predict consensus cleavage signals for furin, three in the case of ADAMTS-5, and one each in the case of ADAMTS-6 and ADAMTS-7. The most carboxyl-terminal furin cleavage site in ADAMTS-5 predicts the processing site for generation of the mature protease. The amino terminus of the mature proteins is predicted to start at the residue immediately following the cleavage sites.

2. **A catalytic domain.** The catalytic domains are very similar to each other and contain eight cysteine residues and a typical reprolysin-type zinc binding signature followed by a "Met-turn". Five cysteine residues are upstream of the zinc binding sequence, while three residues are downstream, an arrangement that is shared with other ADAMTS members. The methionine of the met-turn is not at a constant distance from the zinc-binding signature, but in all three novel proteases, a constant cysteine residue is present in that interval.

3. **A disintegrin-like domain.** The catalytic domain is followed by a domain of 60-90 residues with 35-45% similarity to snake venom disintegrins, but without the canonical cysteine arrangement seen in the latter. This disintegrin-like domain is of comparable length in ADAMTS-5 and ADAMTS-7, it is considerably shorter in ADAMTS-6.

4. **A TS module.** The first TS repeat is very similar in all three novel proteases and very similar to the first TS repeat of other ADAMTSs. It contains the same number of residues (fifty-two) in all three novel proteins.

5. **The cysteine-rich domain.** This TS domain is followed by a

other domains. It shows the least homology of all the domains.

7. **A C-terminal TS module.** The sequence of the second TS module is more variant between the members of the ADAMTS family than the first TS module, despite the conservation of the number and spacing of cysteine residues.

Overall, the predicted mature forms of these proteases show 20-30% similarity to each other and to ADAMTS1-4 although this may be considerably higher or lower for individual domains as described above.

ADAM-TS9 and ADAM-TS10 contain all the domains present in ADAMTS-5 through ADAMTS-8. In addition, ADAMTS-9 and ADAMTS-10 contain the following domains:

A. ADAMTS-9: After the c-terminal TS1 domain which is present in ADAMTS5-8, ADAMTS-9 contains 13 additional and homologous TS11 domains, thus, ADAMTS-9 contains a total of 15 TS1 domains, of which 14 are adjacent to each other in the c-terminal half of the molecule. The 15th TS1 domain from the N-terminus is followed by a unique c-terminal domain which does not possess recognizable domain structure and contains 196 residues including 9 cysteine residues.

B. ADAMTS-10: After the c-terminal TS1 domain which is present in ADAMTS 8, ADAMTS-10 contains 3 additional and homologous TS1 domains, thus, that ADAMTS-10 contains a total of 5 TS1 domains, of which 4 are adjacent to each other in the c-terminal half of the molecule. The 5th TS 1 domain from the N-terminus is followed by an additional 47 amino acid residues including six (6) cysteine residues. These 47 residues have sequence similarity of 30%-40% to

from human and mouse tissues (Clontech, Palo Alto, CA) were hybridized to the [α^{32}P]-dCTP labeled inserts of I.M.A.G.E. clones as per the manufacturer's recommendations followed by autoradiographic exposure for 3-7 days.

In situ hybridization used cryosections of mouse embryos of gestational age 8.5 days and 10.5 days. Embryos were collected with the inclusion of the surrounding uterus and fixed overnight in 4% paraformaldehyde. Sense and anti-sense probes continuously labeled with digoxigenin-UTP (Boehringer-Mannheim, Indianapolis, IN) were transcribed with T7 and T3 RNA polymerases, respectively, using as template a 630 bp EcoRI-SacI fragment from the *Adamts-5* clone 569515 (Fig. 14) cloned into pBluescript SK+ (Stratagene, La Jolla, CA). In situ hybridization was done essentially as previously described in Apte, et al. (1997) J. Biol. Chem. 272:2551-25517, which is specifically incorporated herein by reference, except that sections were predigested with proteinase K (Boehringer-Mannheim, Indianapolis, IN) at a lower, concentration (1 -5 µg/ml) than reported in Apte, et al.. Bound, digoxigenin-labeled probe was detected using an alkaline phosphatase tagged anti-digoxigenin antibody (Boehringer-Mannheim, Indianapolis, IN) and nuclei were counterstained with methyl green.

Specific hybridization of the antisense Adamts-5 probe to sections of 8.5 day-old mouse embryos was obtained, whereas only low background staining was noted with the control sense probe. Staining was uniform throughout the 8.5 day old embryos. In addition, there was labeling of mRNA in trophoblastic cells lining the uterine cavity

embryos. labeling was widespread but less intense compared to the

day-old embryo. Labeled cells were seen in mesenchyme and somites as well as in the neural tube and developing hindgut. Northern analysis also indicated that mRNA encoding ADAMTS-5 was present in human placenta but was barely detectable in adult lung, heart, brain, liver, skeletal muscle, kidney and pancreas.

Northern analysis showed undetectable expression of *Adamts-6* during mouse embryo development. Northern analysis indicated that mRNA encoding ADAMTS-6 was present in human placenta but was barely detectable in adult lung, heart, brain, liver, skeletal muscle, kidney and pancreas. *Adamts-7* was expressed at low levels throughout mouse development. In adult human tissues examined with human cDNA probes, ADAMTS-7 mRNA was found in all tissues examined, i.e. in lung, heart, brain, liver, skeletal muscle, kidney, pancreas and placenta. The sizes of the mRNA species recognized by the probes varied. ADAMTS-5 mRNA was approximately 10 kbp in size in human tissue. The most prominent *Adamts-5* species was estimated at 7.5 kbp together with additional bands at 10 kbp and 4.5 kbp. The lone mRNA species detected by ADAMTS-6 probe was approximately 8.5 kbp, whereas the most common mRNA species detected by ADAMTS-7 probe 5 was 5 kbp in size with an additional species seen at 7 kbp in skeletal muscle.

In mouse, ADAMTS-8 is expressed during fetal development (days 7, 11, 15, 17) and in adult mouse lung and heart with an mRNA size of approximately 3.8 kbp. In adult human tissue, ADAMTS-8 is expressed in lung and brain but not in heart, muscle, kidney, colon or thymus. The mRNA size is 3.8 kbp.

ADAMTS-9 is expressed in lung, ovary placenta, heart, brain,

1 alternatively spliced is short form of ADAMTS

ADAMTS-10 is expressed in thymus, prostate, testis, ovary, small intestine, colon, peripheral blood leukocytes, heart, brain, placenta, lung, liver, muscle, kidney and pancreas, as well as in many cell lines such as A549, HeLa and K562. There are two transcripts of 5 kb and 8kb present in all tissues.

Example 7: ADAMTS-R1

The nucleotide sequence of a cDNA encoding a full-length ADAMTS-R1 protein was obtained using IMAGE clone 752797 which encodes EST AA, and a human fetal brain cDNA from Clontech. RACE was performed as described above in Example 1. The nucleotide sequence, SEQ ID NO:21, of the ADAMTS-R1 cDNA and the predicted amino acid sequence, SEQ ID NO:22, of the ADAMTS-R1 protein encoded by such DNA is shown in Fig. 11.

The predicted Mr of the full-length, unprocessed ADAMTS-R1 protein is 58358.20 daltons. The domain organization of the ADAMTS-10 protein is shown in Fig. 15. In contrast to the ADAMTS-N proteins of examples 1-6, ADAMTS-R1 protein does not have a pro-metalloprotease or disintegrin-like domain or a consensus cleavage signal for furin. ADAMTS-R1 has a signal(pre) peptide which is followed by a first TS module and a conserved CRD sequence which contains 10 conserved cysteines. The spacer domain of ADAMTS-R1 is 119 amino acids in length and is followed by 3 additional TS modules and a short sequence of 33 amino acids. The ADAMTS-R1 protein contains one potential glycosylation sites which is in the spacer domain. ADAMTS-R1 bears 30-40% sequence identity to ADAMTS1 and ADAMTS4 in the related domains. ADAMTS-R1 mRNA is present in human heart, brain, kidney, muscle, lung, placenta, testis, ovary, colon,

Although certain embodiments of this invention have been shown and described, various adaptations and modifications can be made without departing from the scope of the invention as defined in the appended claims.

CLAIMS

1. An isolated mammalian protein selected from the group consisting of an ADAMTS-5 protein an ADAMTS-6 protein, an ADAMTS-7 protein, an ADAMTS-8 protein, an ADAMTS-9 protein, an ADAMTS-10 protein, and an ADAMTS-R1 protein.

2. The isolated mammalian protein of claim 1 wherein said protein comprises an amino acid sequence which is at least 95% identical to a sequence selected from the group consisting of: amino acid 262 through amino acid 930 of SEQ ID NO:2; amino acid 1 through amino acid 518 of SEQ ID NO:4; amino acid 245 through amino acid 860 of SEQ ID NO:6; amino acid 233 through amino acid 997 of SEQ ID NO:8; amino acid 229 through amino acid 905 of SEQ ID NO:10; amino acid 1 through amino acid 245 of SEQ ID NO:12; amino acid 236 through amino acid 1882 of SEQ ID NO:14; amino acid 1 through amino acid 874 of SEQ ID NO:16; amino acid 212 through amino acid 1081 of SEQ ID NO:18; amino acid 1 through amino acid 450 of SEQ ID NO:20; and amino acid 1 through amino acid 547 of SEQ ID NO:22.

3. The isolated protein of claim 2 wherein said amino acid sequence further comprises a prepropeptide sequence at the amino terminus thereof.

4. The isolated protein of claim 1 wherein said protein is a human ADAMTS-5 protein or a mouse ADAMTS-5 protein.

5. The isolated protein of claim 1 wherein said protein is a human ADAMTS-6 protein.

6. The isolated protein of claim 1 wherein said protein is a human

...

8. The isolated protein of claim 1 wherein said protein is a human

ADAMTS-9 or a mouse ADAMTS-9 protein.

9. The isolated protein of claim 1 wherein said protein is a human ADAMTS-10 or a mouse ADAMTS-10 protein.

10. The isolated protein of claim 1 wherein said protein is a human ADAMTS-R1 protein.

11. An isolated polynucleotide comprising a sequence which encodes a mammalian protein selected from the group consisting of an ADAMTS-5 protein, an ADAMTS-6 protein, an ADAMTS-7 protein, an ADAMTS-8 protein, an ADAMTS-9 protein, an ADAMTS 10 protein, and an ADAMTS-R1 protein.

12. The isolated polynucleotide of claim 11 wherein said protein comprises an amino acid sequence which is at least 95% identical to a sequence selected from the group consisting of: amino acid 262 through amino acid 930 of SEQ ID NO:2; amino acid 1 through amino acid 518 of SEQ ID NO:4; amino acid 245 through amino acid 860 of SEQ ID NO:6; amino acid 233 through amino acid 997 of SEQ ID NO:8; amino acid 229 through amino acid 905 of SEQ ID NO:10; amino acid 1 through amino acid 245 of SEQ ID NO:12; amino acid 236 through amino acid 1882 of SEQ ID NO:14; amino acid 1 through amino acid 874 of SEQ ID NO:16; amino acid 212 through amino acid 1081 of SEQ ID NO:18; amino acid 1 through amino acid 450 of SEQ ID NO:20, and amino acid 1 through amino acid 847 of SEQ ID NO:22.

13. The isolated polynucleotide of claim 11 wherein said nucleotide sequence encodes a protein having a signal sequence at the amino terminus thereof.

[illegible] NO:1 of an allelic variant thereof; nucleotide 1 through

nucleotide 1519 of SEQ ID NO:3 or an allelic variant thereof; nucleotide 754 through nucleotide 2602 of SEQ ID NO:5 or an allelic variant thereof; nucleotide 708 through nucleotide 3003 of SEQ ID NO:7 or an allelic variant thereof; nucleotide 962 through nucleotide 2992 of SEQ ID NO:9 or an allelic variant thereof; nucleotide 1 through nucleotide 739 of SEQ ID NO:11 or an allelic variant thereof; nucleotide 708 through nucleotide 5648 of SEQ ID NO:13 or an allelic variant thereof; nucleotide 1 through nucleotide 2625 of SEQ ID NO:15 or an allelic variant thereof; nucleotide 634 through nucleotide 3243 of SEQ ID NO:17 or an allelic variant thereof; nucleotide 1 through nucleotide 1642 of SEQ ID NO:19 or an allelic variant thereof; and nucleotide 51 through nucleotide 1625 of SEQ ID NO:21 or an allelic variant thereof.

15. The isolated polynucleotide of claim 11 wherein said polynucleotide hybridizes under stringent conditions to a nucleic acid molecule comprising a sequence complementary to the protein encoding sequence of SEQ ID NO:1; SEQ ID NO:3; SEQ ID NO:5; SEQ ID NO:7; SEQ ID NO:9; SEQ ID NO:11; SEQ ID NO:13; SEQ ID NO:15; SEQ ID NO:17; SEQ ID NO:19; or SEQ ID NO:21.

16. An isolated polynucleotide having a sequence which is complementary to the protein encoding sequence of the polynucleotide of claim 11.

17. An expression vector comprising a polynucleotide of claim 11.

18. A host cell transformed or transfected with an expression vector of claim 17.

protein; and

(b) recovering said ADAMTS-N protein or said ADAMTS-R1 protein from the host cell culture.

20. An antibody that binds to a protein selected from the group consisting of an ADAMTS-5 protein, an ADAMTS-6 protein, an ADAMTS-7 protein, an ADAMTS-8 protein, an ADAMTS-9 protein, an ADAMTS-10 protein and an ADAMTS-R1 protein.

21. An oligopeptide for producing an antibody that binds to an ADAMTS N protein or an ADAMTS-R1 protein wherein said oligopeptide has a sequence selected from the group consisting of:

a) SVSIERFVETLVVADK, SEQ ID NO:23;

b) EVAEAANFLALRSEDPDKY, SEQ ID NO:24;

c) VKEDVENPKAVVDGDWGP, SEQ ID NO:25;

d) QHPFQNEDYRPRSASPSRTH, SEQ ID NO:26;

e) PQNCKEVKRLKGASEDGEYF, SEQ ID NO:27;

f) QELEEGAAVSEEPS, SEQ ID NO:28;

g) YYPENIKPKPKLQE; SEQ ID NO:29;

h) HIKVRQFKAKDQTRF; and

i) CEAKNGYQSDAKGVKTFVEWVPKYAG, SEQ ID NO:30.

Fig. 1

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Fig. 2

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Fig. 3

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Fig. 3 (con't)

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Fig. 4

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Fig. 5A

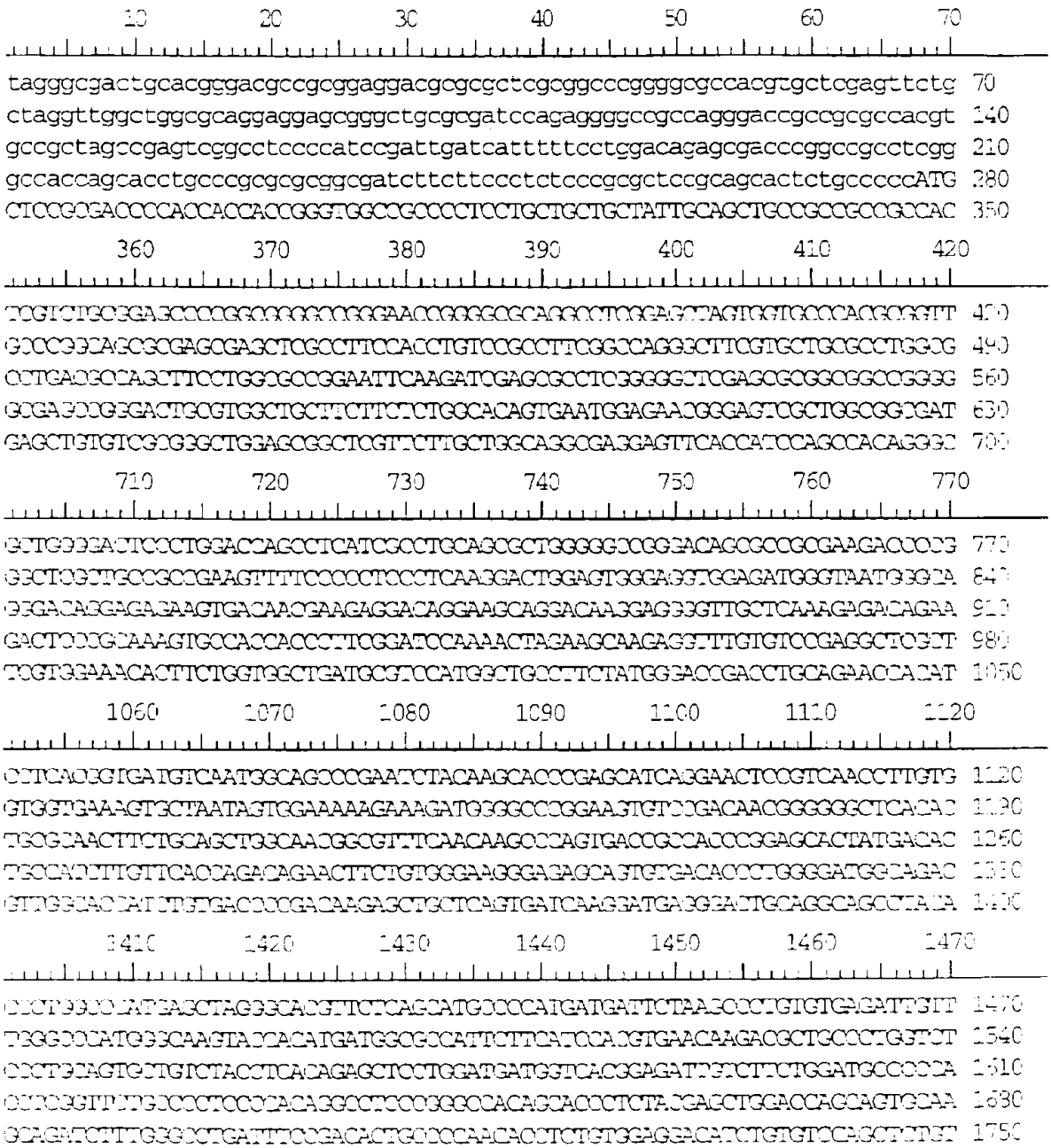
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Fig. 5A (con't)

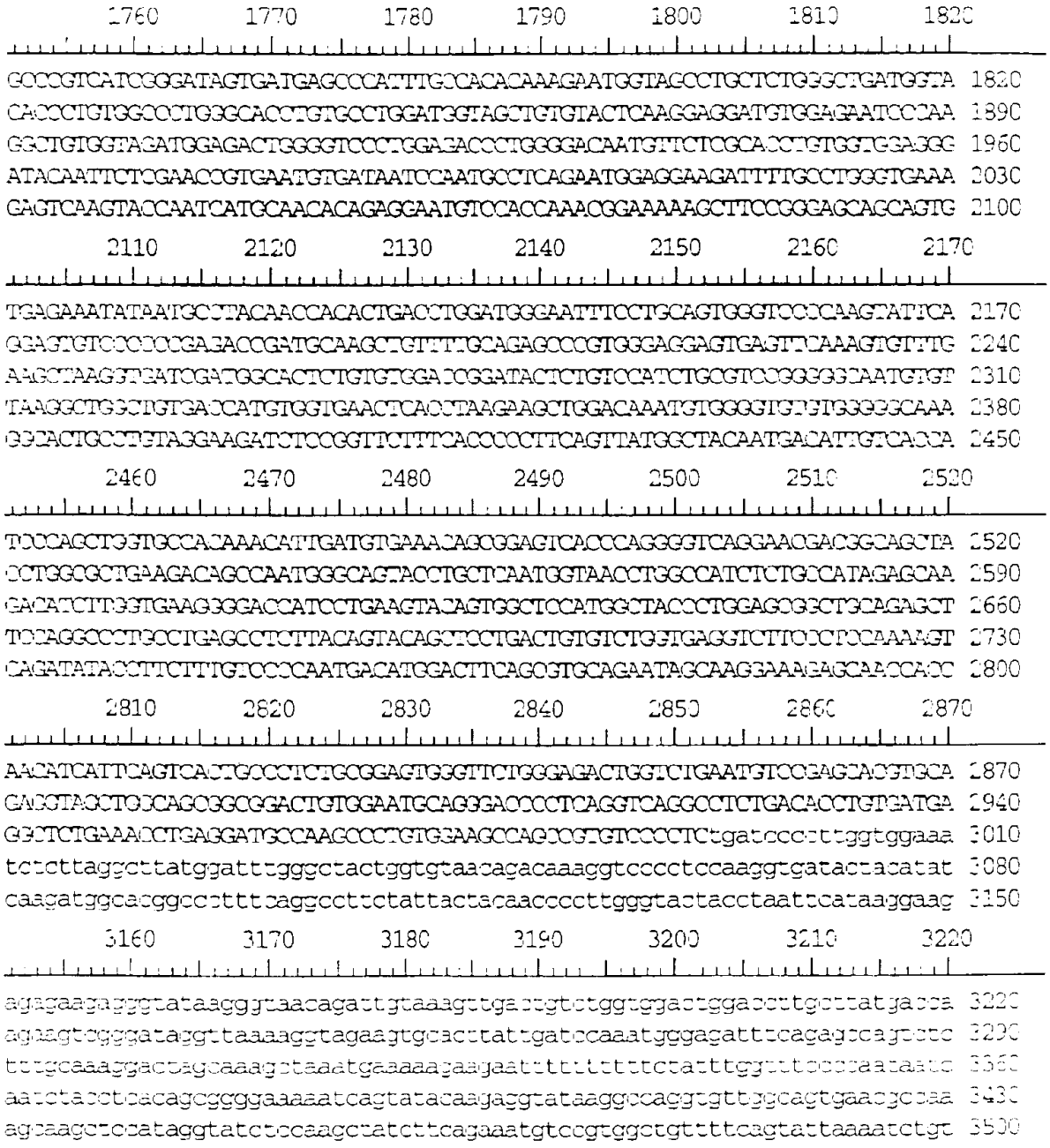
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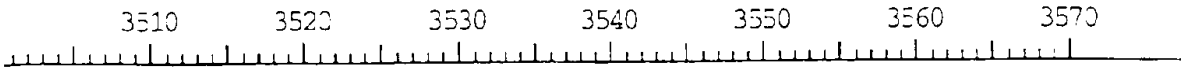
Fig. 5A (con't)



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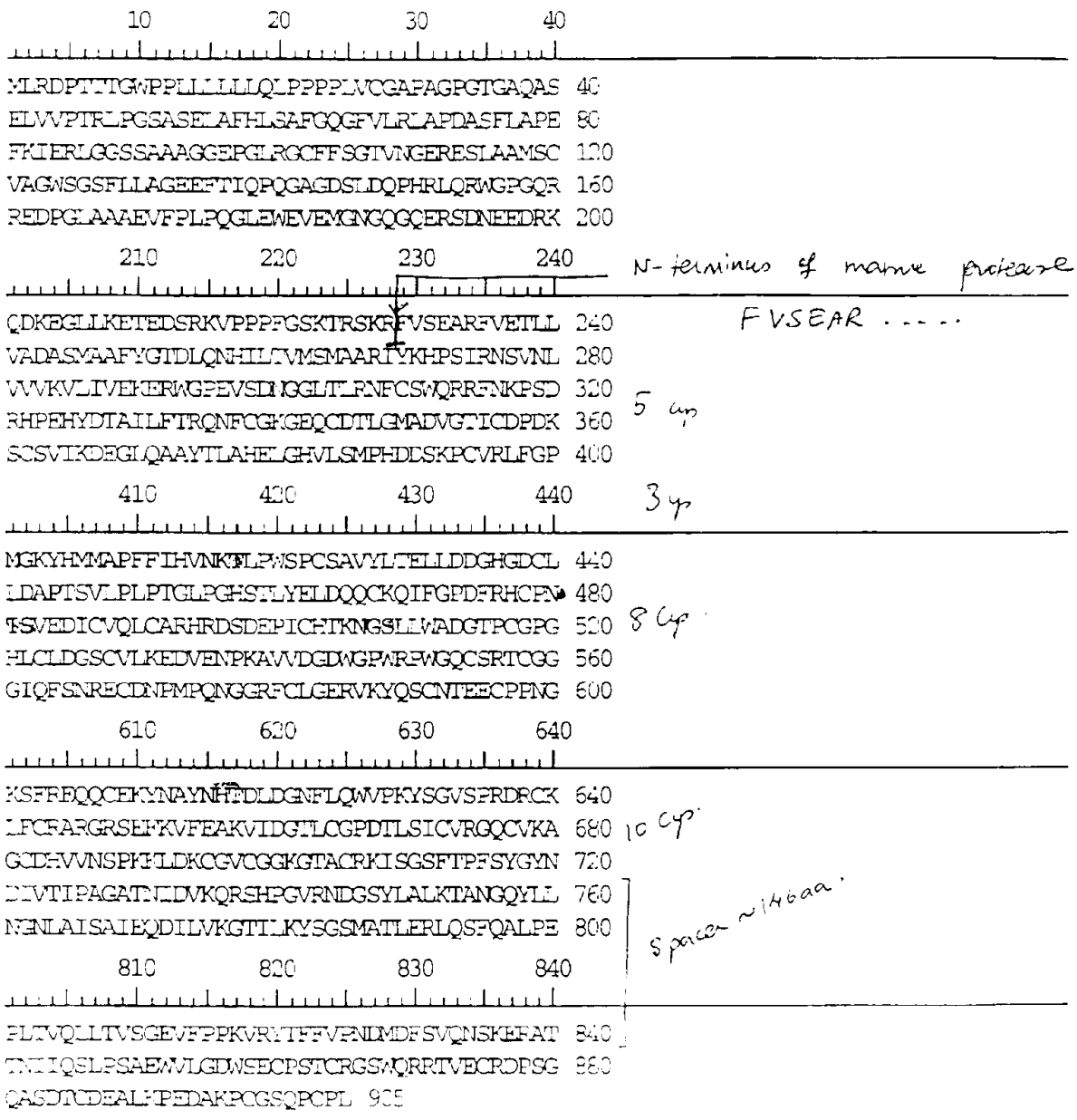
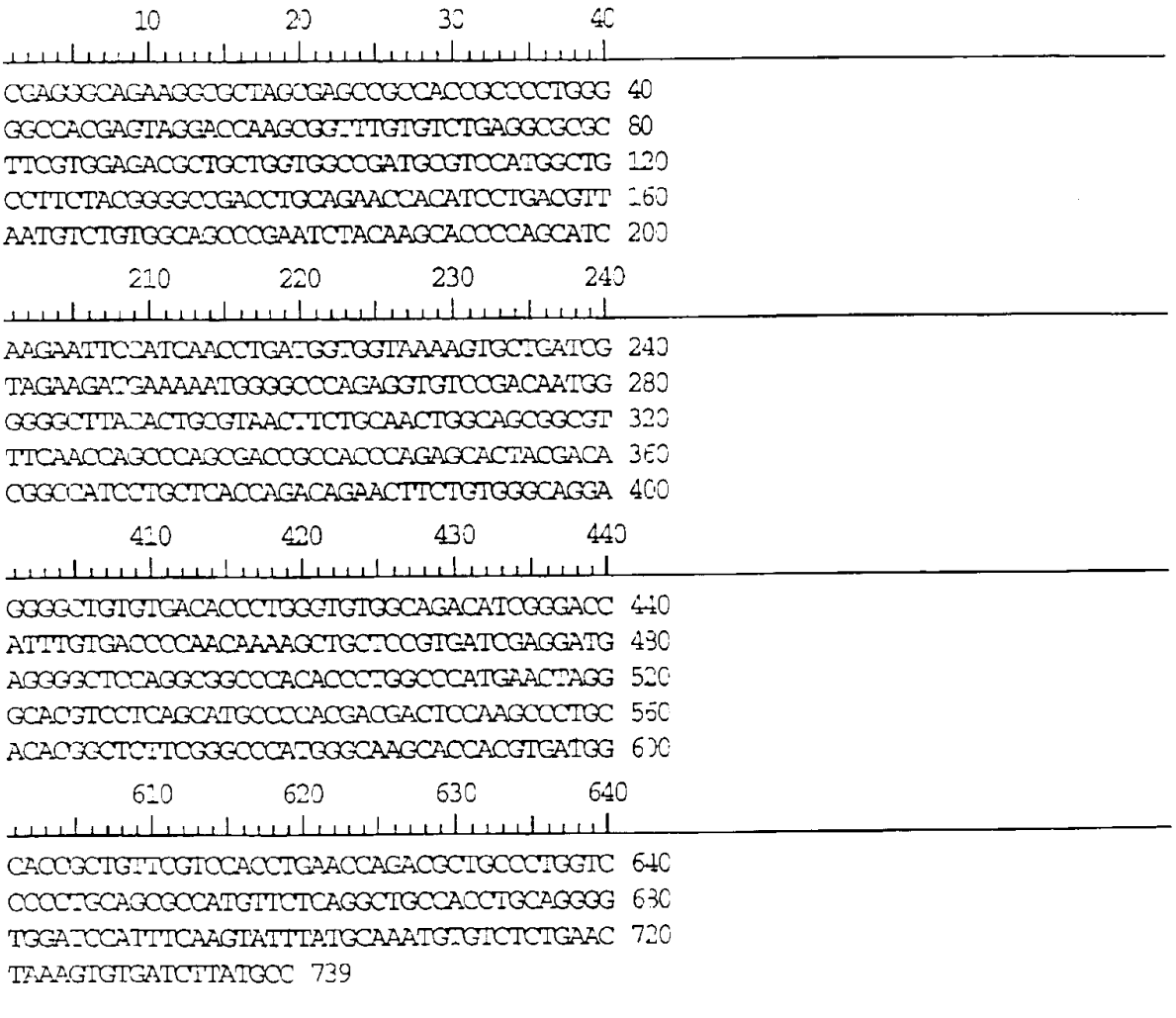


Fig. 6A

CATALYTIC DOMAIN, ADAM TS-8 (HUMAN)



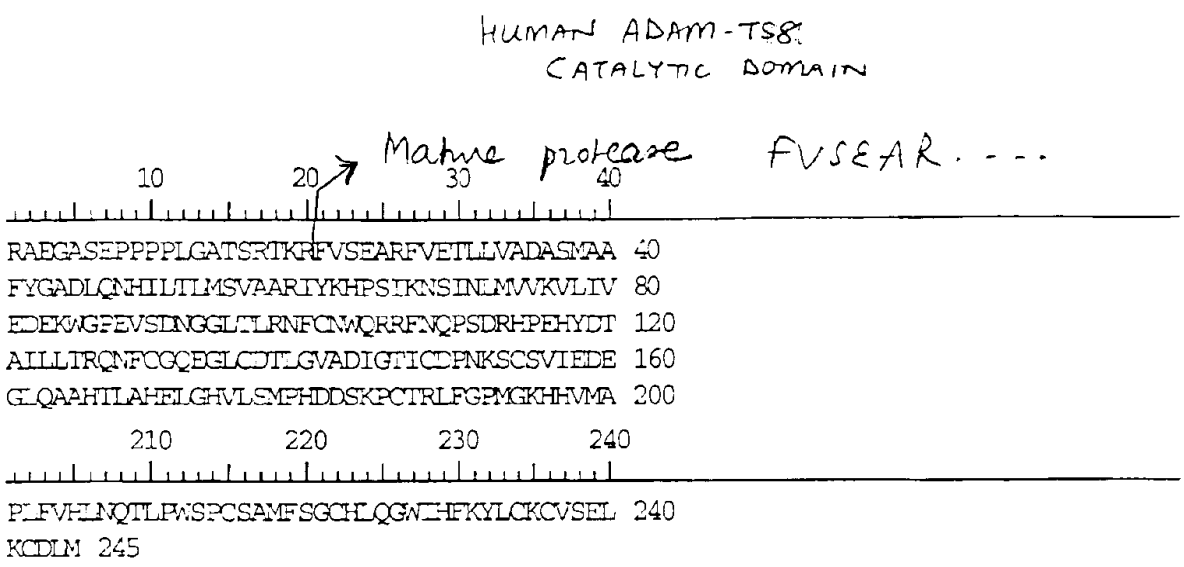

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Fig. 6B

Fig. 7A

Human HDAm-T19

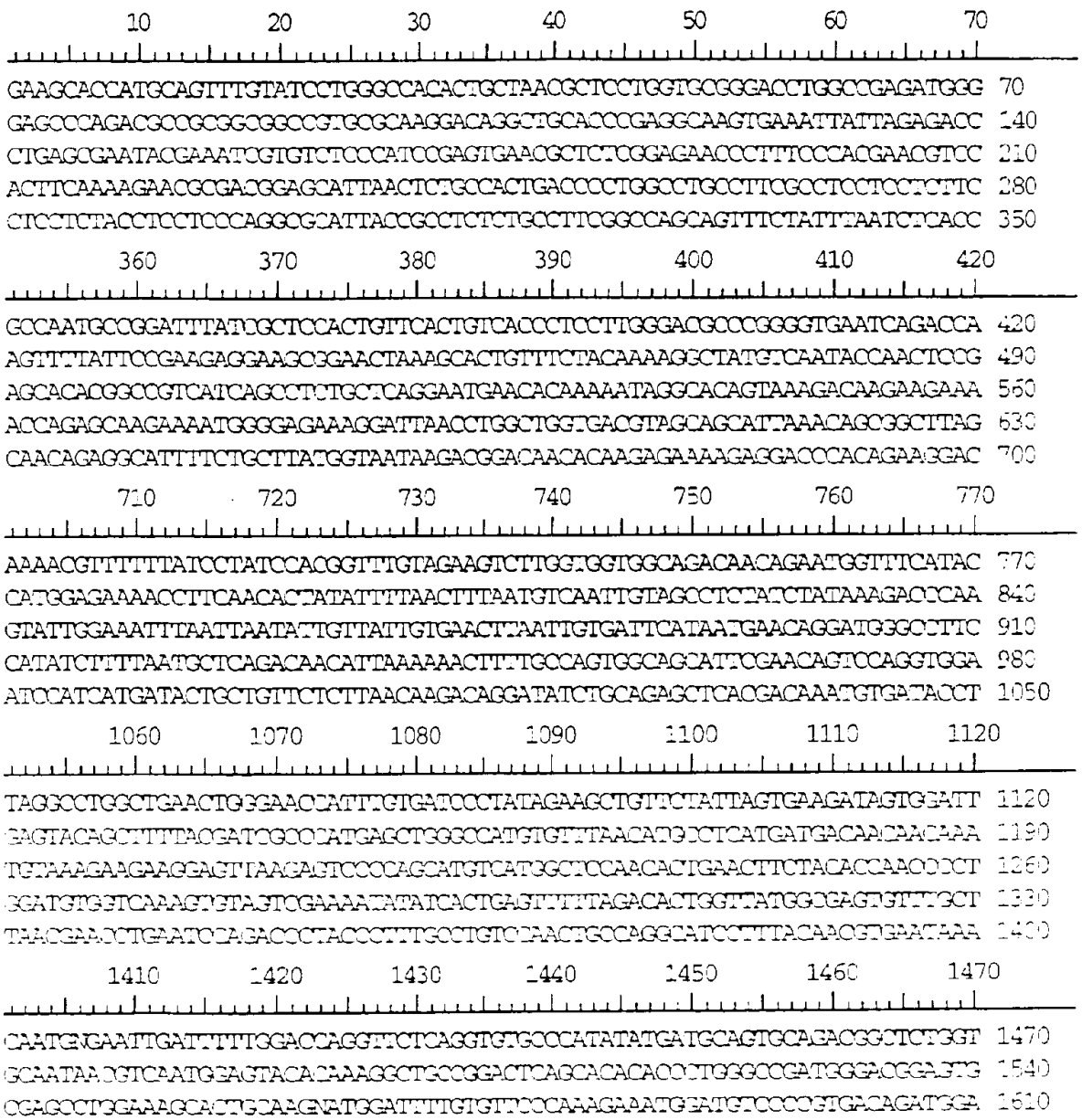
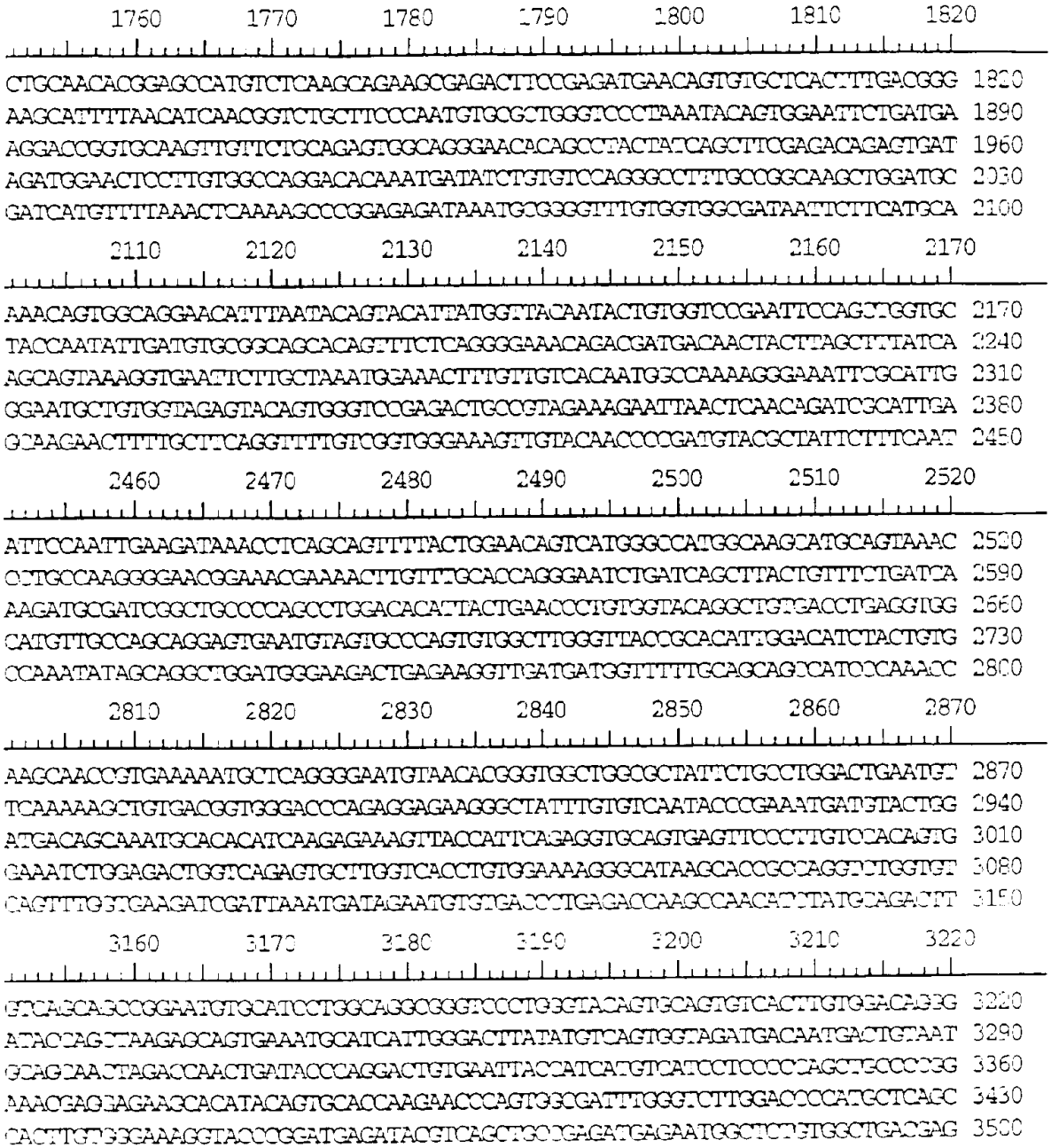


Fig. 7A (con't)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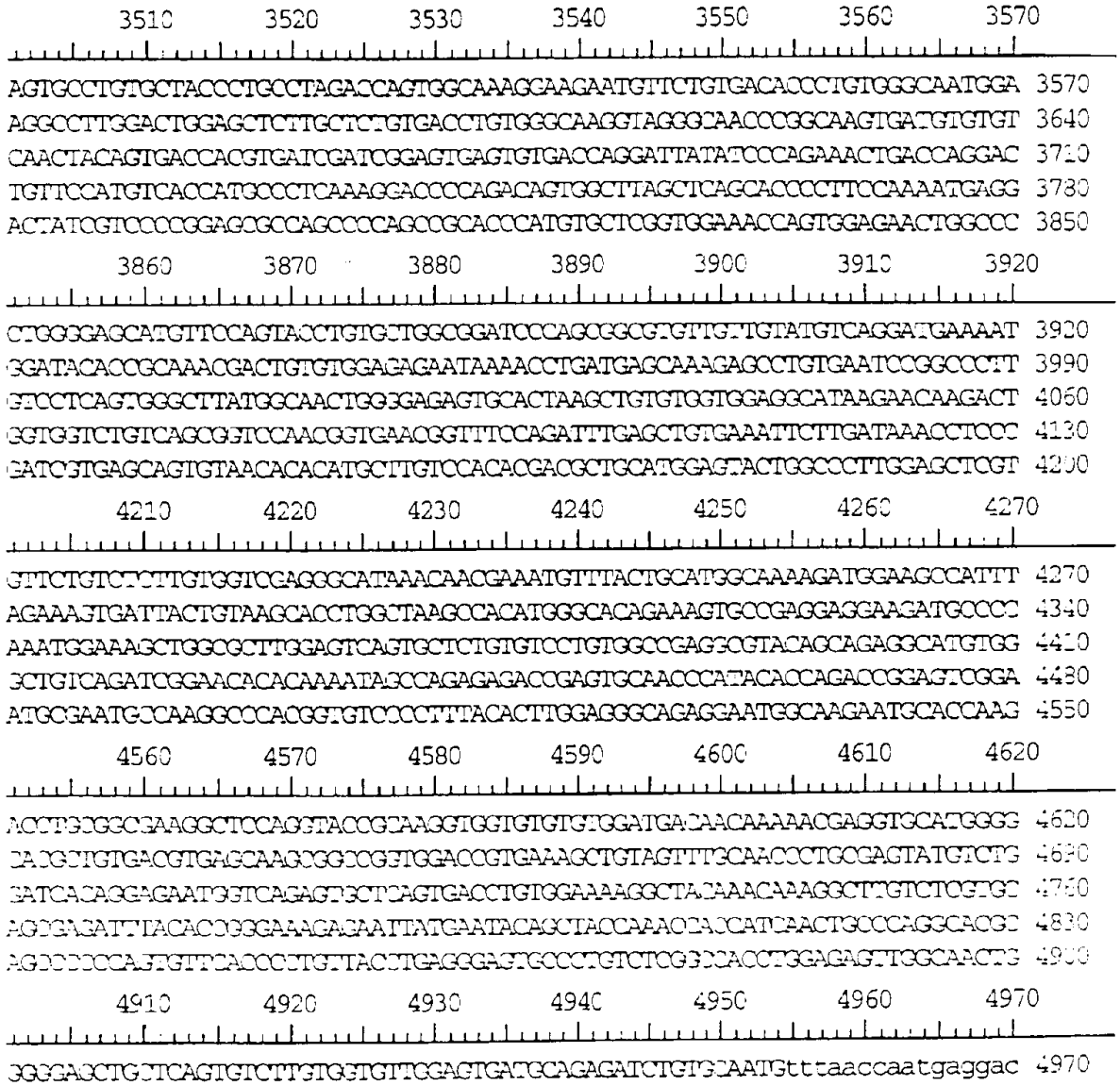
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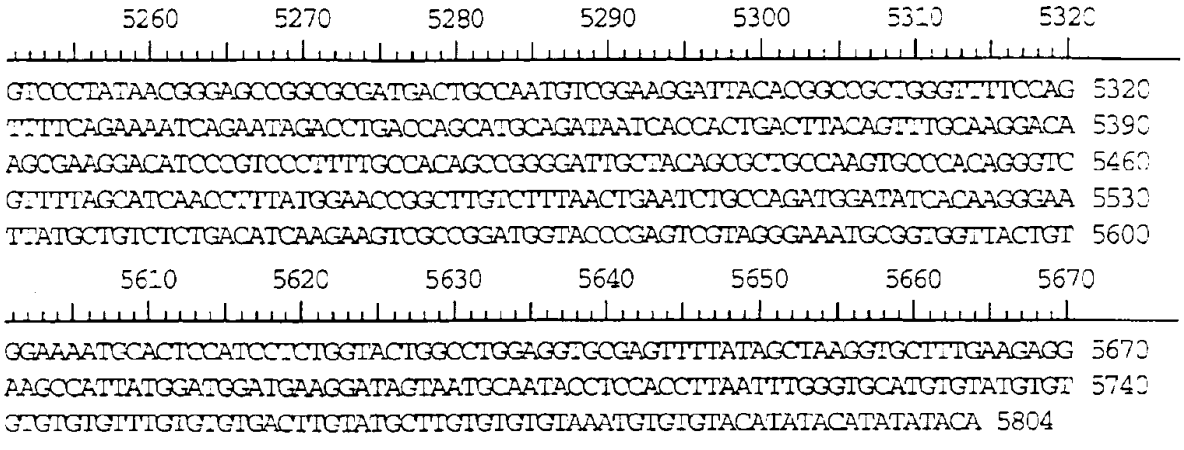
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Fig. 7B

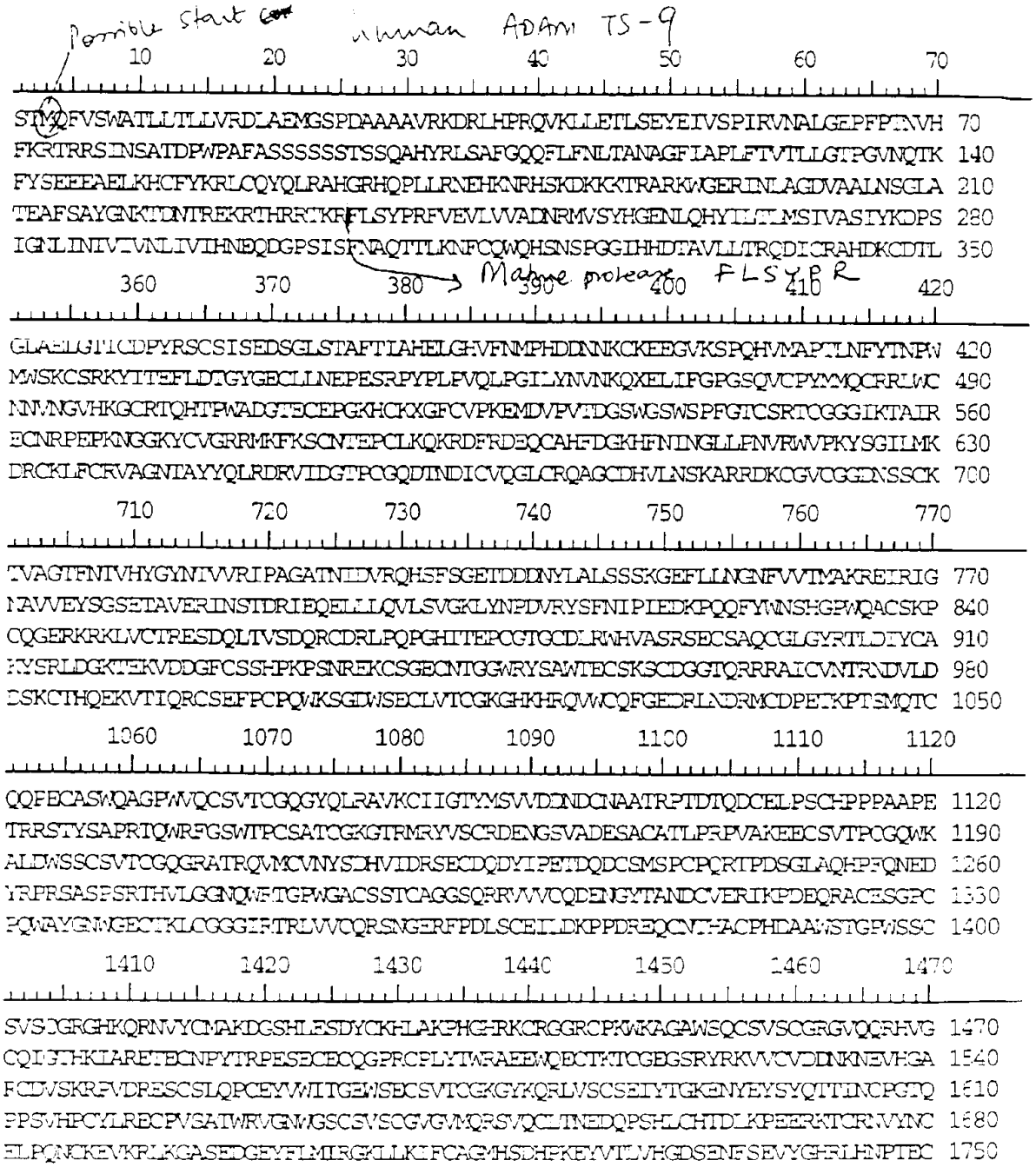

Fig. 7B (con't)

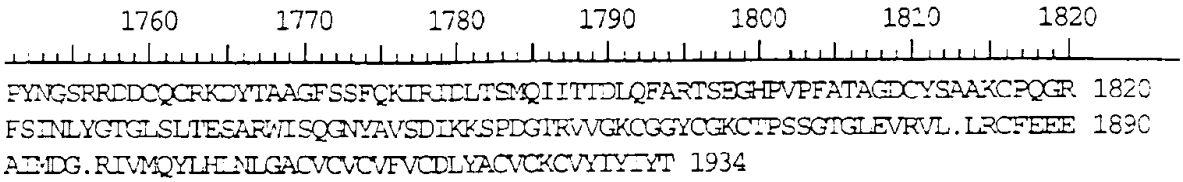
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Fig. 8

ORF=2 protein

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maine ADAMTS9
FLSYPRF...
Mouse ADAM-TS9
partial segment
(see figure)

Created: Saturday, April 10, 1999 11:40 AM

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Fig. 8 (con't)

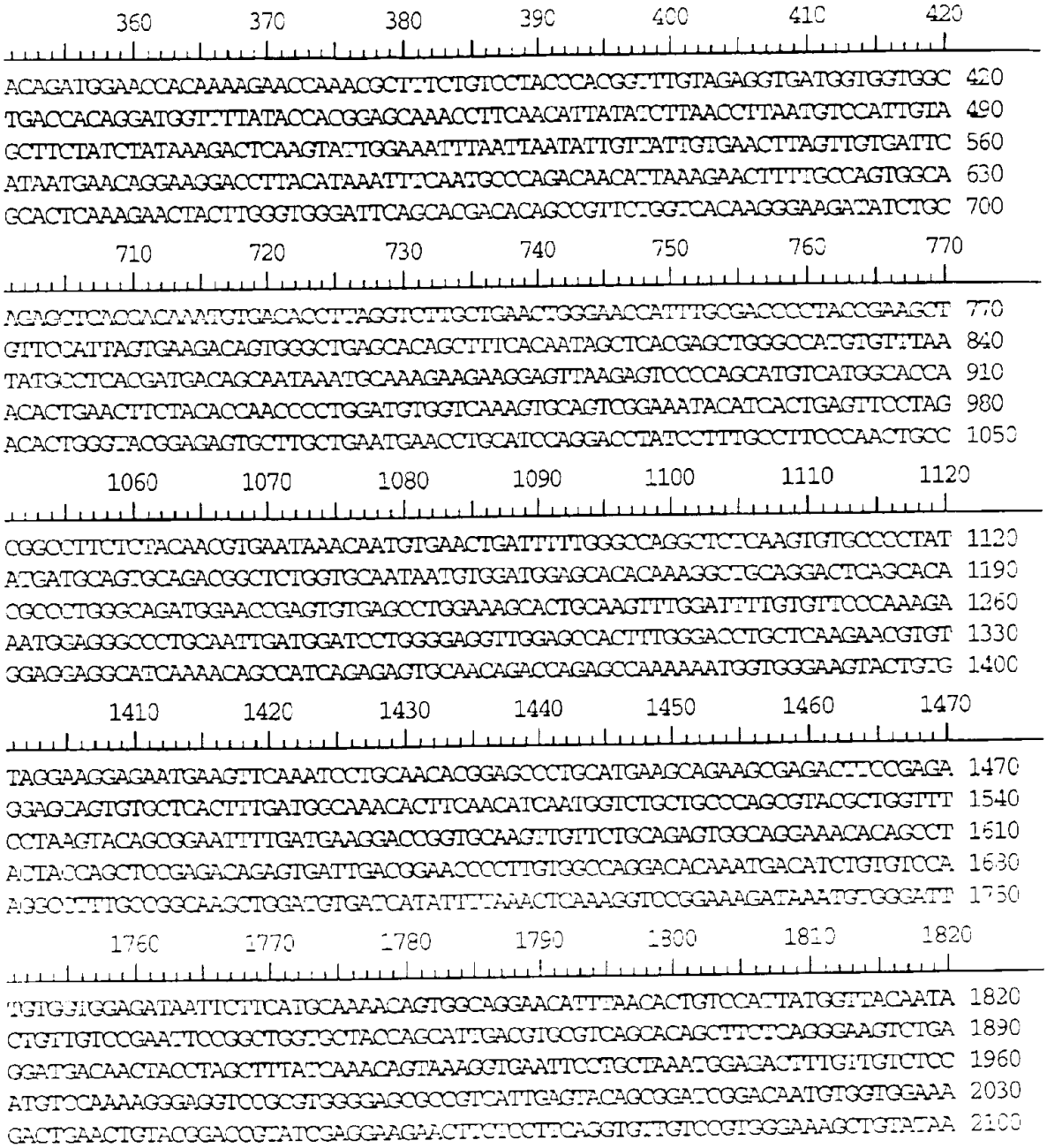
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Fig. 8 (con't)

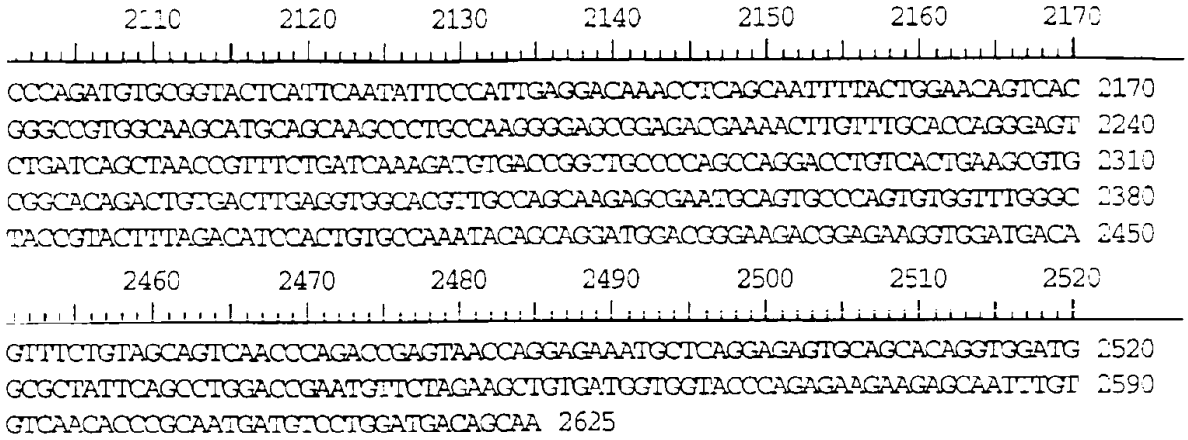
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Fig. 9A

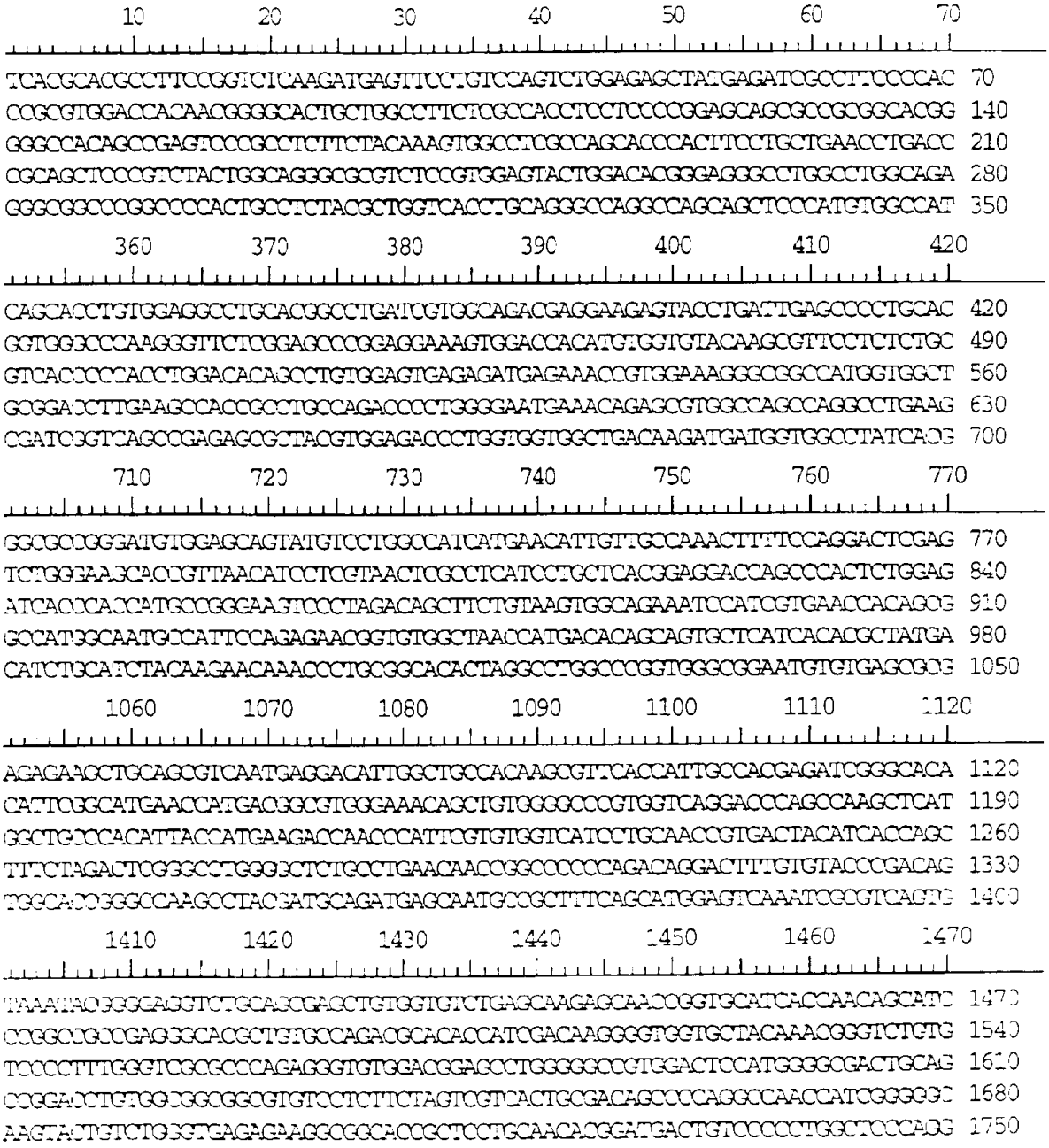
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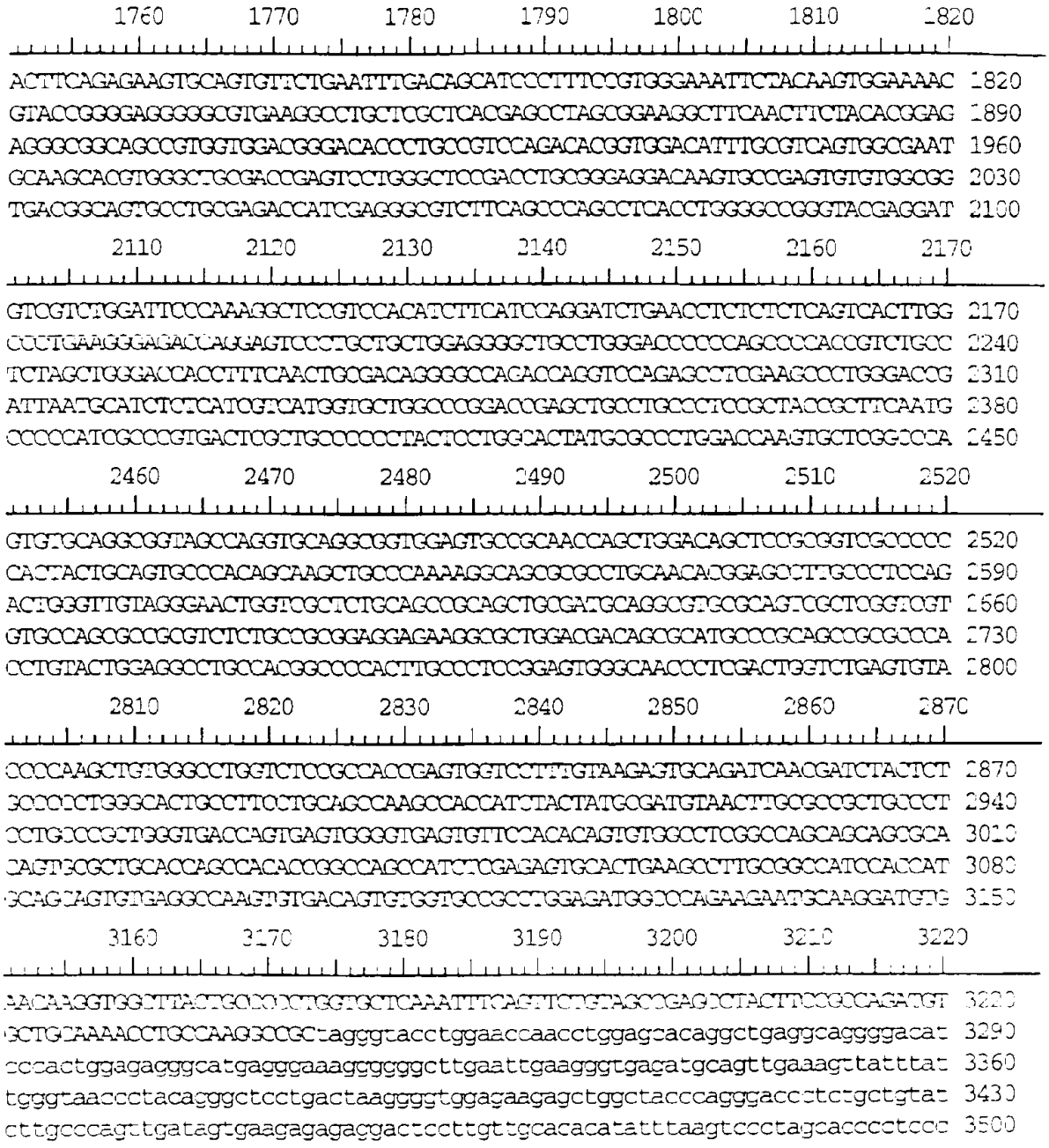
Fig. 9A (con't)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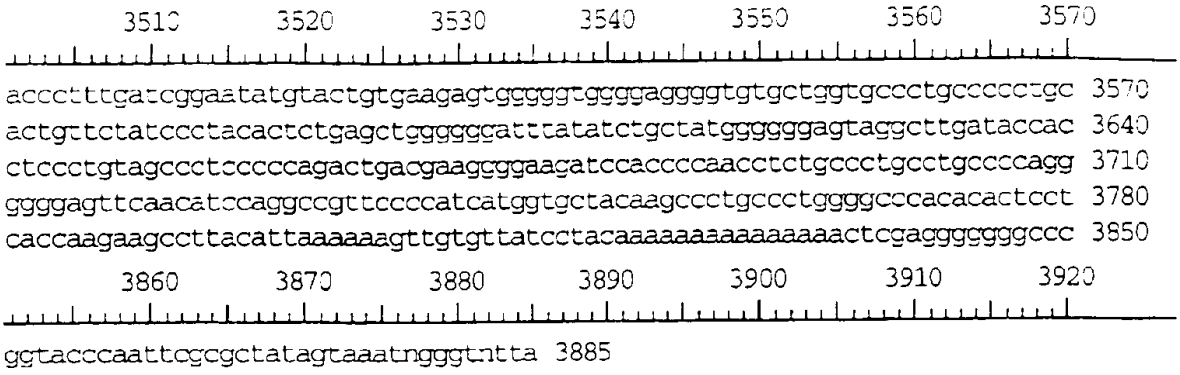
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Fig. 9B

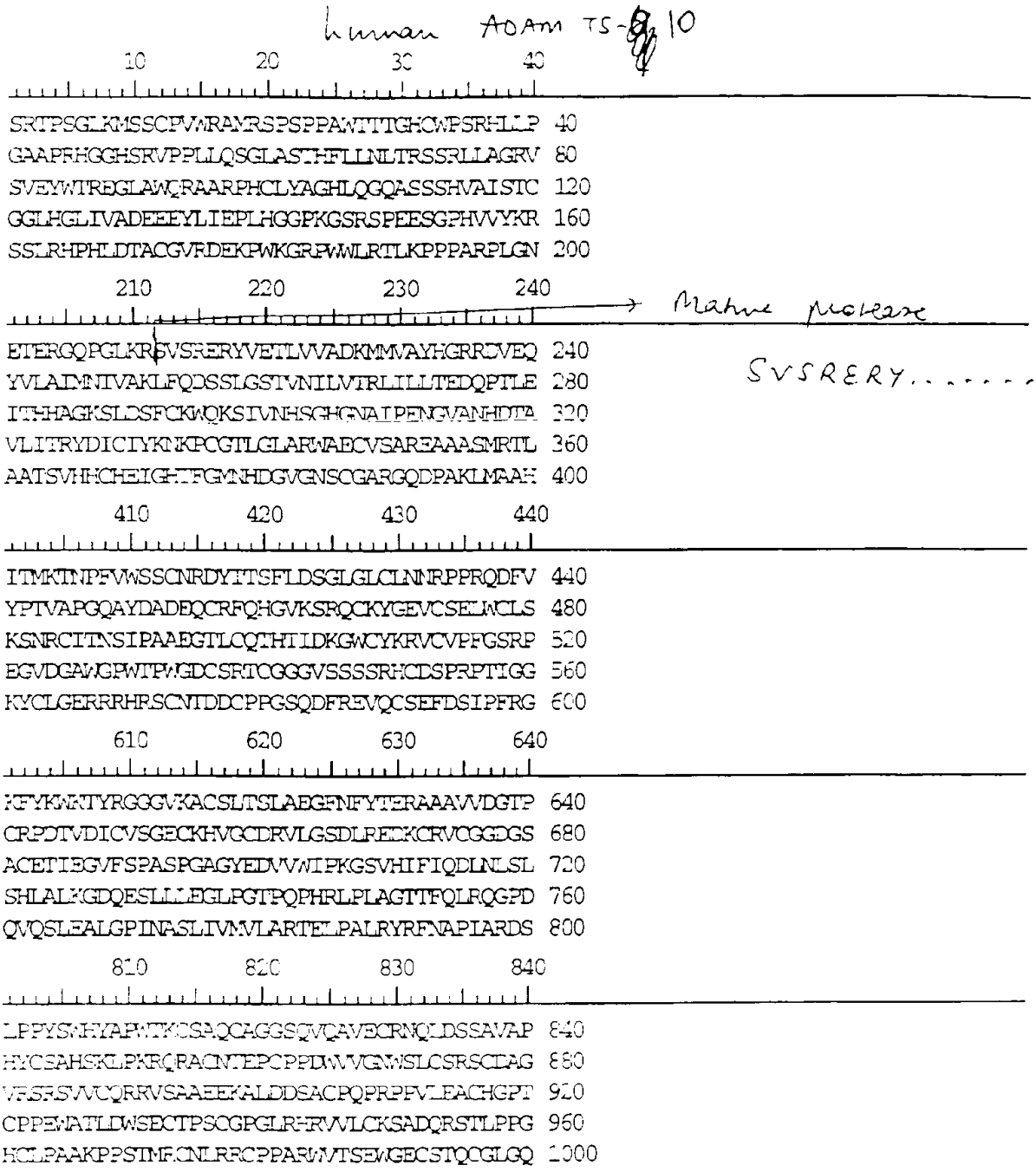
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Fig. 9B (con't)

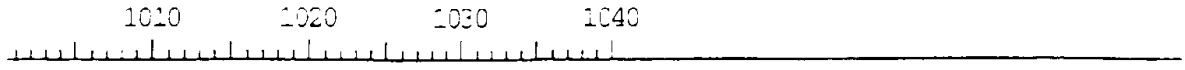

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Fig. 10A

partial sequence of mouse ADAM TS-10 (see figure)

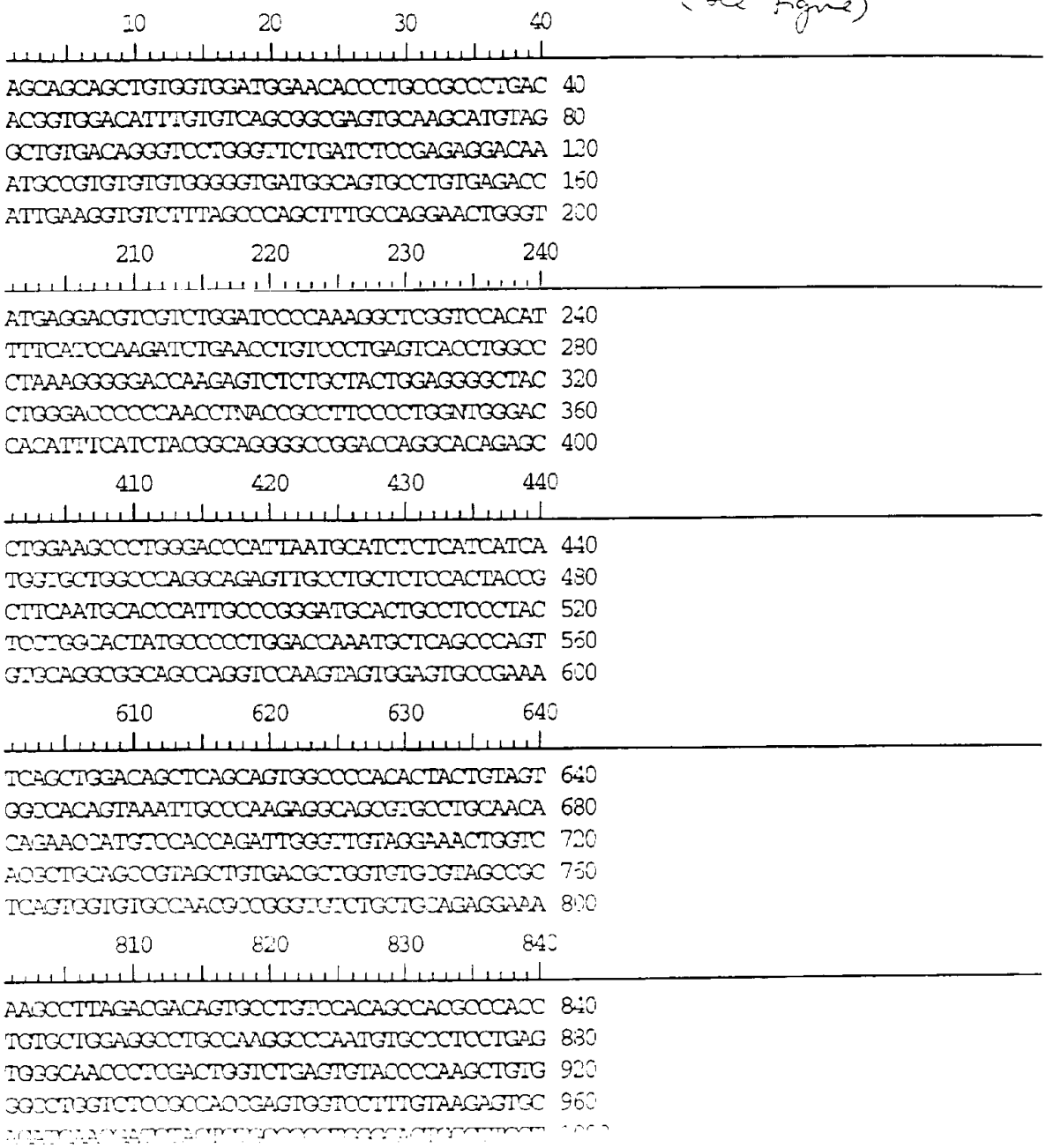

Fig. 10A (con't)

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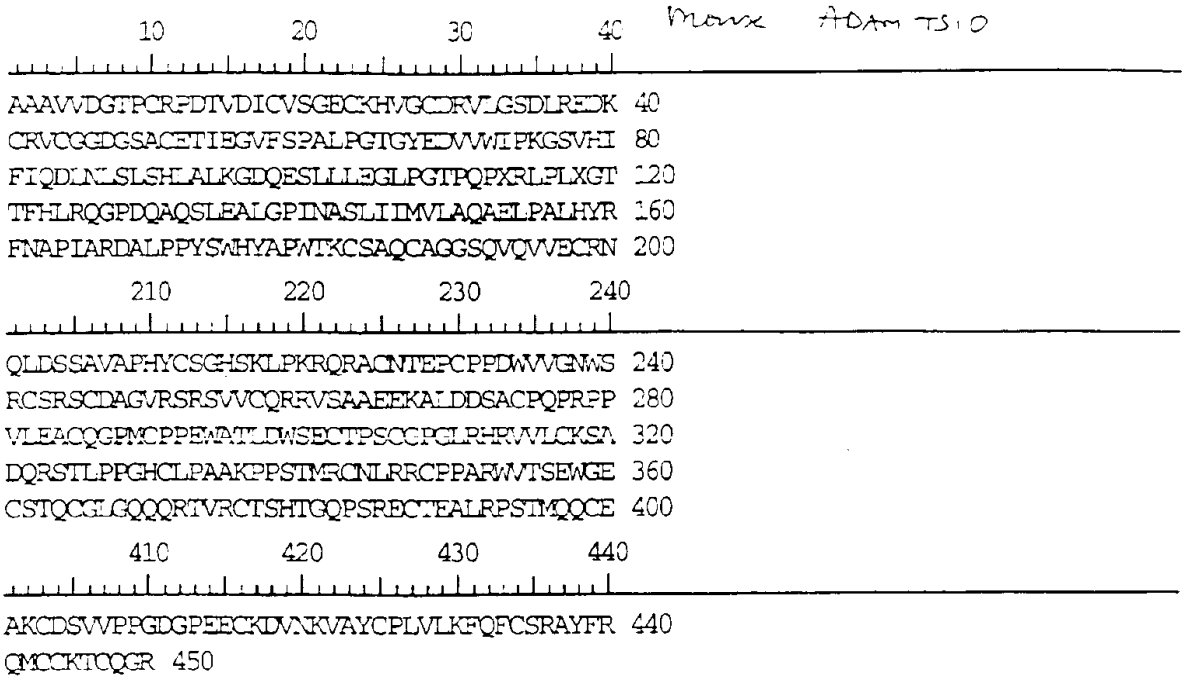
Fig. 10B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QMCCKTCQGR 450

Fig. 11A

Ligated 459225+482392 with Sac I(168)&Eco RI(or Not I)
Cloning site:5';Eco RI 3';Not I Vector; PI7T3 pac.

You can put this construct to pcDNA3.1(+) for transfection
5'-UTR is 50bp &3'-UTR is 175bp

210-215; in 482392 it's TCCTAC(SY).

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Fig. 11A (con't)

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CAAGATTCGTAACTCGGGCTCCGCTGACAGTACAGTCCAG 920
TTCATCTTCTATCAACCCATCATCCACCGATGGAGGGAGA 960
CGGATTTCTTTCCTTGCTCAGCAACCTGTGGAGGAGGTTA 1000
        1010       1020       1030       1040
TCAGCTGACATCGGCTGAGTGCTACGATCTGAGGAGCAAC 1040
CGTGTGGTTGCTGACCAATACTGTCACTATTACCCAGAGA 1080
ACATCAAACCCAAACCCAAGCTTCAGGAGTGCAACTTGGA 1120
TCCTTGTCCAGCCAGTGACGGATACAAGCAGATCATGCCT 1160
TATGACCTCTACCATCCCCTTCCTCGGTGGGAGGCCACCC 1200
        1210       1220       1230       1240
CATGGACCGCGTGCTCCTCCTCGTGTGGGGGGGCATCCA 1240
GAGCCGGGCAGTTCCTGTGTGGAGGAGGACATCCAGGGG 1280
CATGTCACTTCAGTGGAAGAGTGGAAATGCATGTACACCC 1320
CTAAGATGCCCATCGCGCAGCCCTGCAACATTTTTGACTG 1360
CCCTAAATGGCTGGCACAGGAGTGGTCTCCGTGCACAGTG 1400
        1410       1420       1430       1440
ACGTGTGGCCAGGGCCTCAGATACCGTGTGGTCCTCTGCA 1440
TCGACCATCGAGGAATGCACACAGGAGGCTGTAGCCCAAA 1480
AACAAAGCCCCACATAAAAGAGGAATGCATCGTACCCACT 1520
CCCTGCTATAAACCCAAAGAGAAACTTCCAGTCGAGGCCA 1560
AGTTGCCATGGTTCAAACAAGCTCAAGAGCTAGAAGAAGG 1600
        1610       1620       1630       1640
AGCTGCTGTGTCAGAGGAGCCCTCGTAAgttgtaaaagca 1640
cagattgtttctatatttgaaacttttgtttaaagaaagca 1680
gtgtctcactggttgtagctttcatgggttctgaactaag 1720
tgtaatcatctcaccaaagcttttttggctctcaaattaaa 1760
gattgattagtttcaaaaaaaaaaaaaaaaaaagatgcggc 1800
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g. 11A (con't)

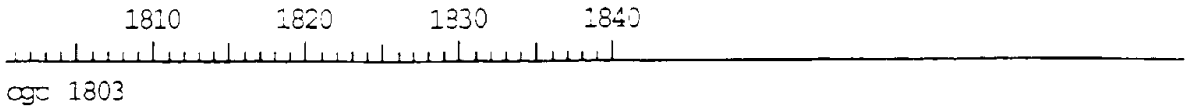


Fig. 11B

---	Asp(D)	30	# cua	Leu(L)	3	# uca	Ser(S)	6	# guu	Val(V)	6
ugc	Cys(C)	26	# cuc	Leu(L)	11	# ucc	Ser(S)	10	# ---	Val(V)	29
ugu	Cys(C)	10	# cug	Leu(L)	14	# ucg	Ser(S)	5	# nnn	???(X)	0
---	Cys(C)	36	# cuu	Leu(L)	6	# ucu	Ser(S)	5	# TOTAL		526
caa	Gln(Q)	7	# uua	Leu(L)	4	# ---	Ser(S)	43	#		

Created: Wednesday, May 5, 1999 10:19 AM

Ligated 459225+482392 with Sac I(168)&Eco RI(or Not I)
Cloning site:5';Eco RI 3';Not I Vector; PT7T3 pac.

...

human ADAM-TSR1

Adam-TS related protein-1.

```
          10        20        30        40
          |         |         |         |
MECCRRATPGTLLLFLAFLLLSSRTARSEEDRDGLWDAWG 40
PWSECSRTCGGGAANSLPRCLSSKSCEGRNIRYRTCSNVD 80
CPPEAGDFRAQQCSAHNDVKHHGQFYEWLPVSNDFDNPCS 120
LKCQAKGITLVVELAPKVLDGTRCYTESLDMCISGLCQIV 160
GCDHQLGSTVKEDNCGVCNGDGSTCRLVRGQYKSQLSATK 200
          210       220       230       240
          |         |         |         |
SDDTVVAIPYGSRHIRLVLKGPDHLYLETKTLQGTKGENS 240
LSSTGTFLVDNSSVDFQIFPDKEILRMAGPLTADFIVKIR 280
NSGSADSTVQFIFYQPIIHRWRETDFFPCSATCGGGYQLT 320
SAECYDLRSNRVVADQYCHYYPENIKPKPKLQECNLDPCP 360
ASDGYKQIMPYDLYHPLPFWEATPWTACSSSCGGGIQSRA 400
          410       420       430       440
          |         |         |         |
VSCVEEDIQGHVTSVEENKCMYTPKMPIAQPCNIFDCPKW 440
LAQEWSPCTVTCGQGLRYRVVLCIDHRGMHIGCCSPKTKP 480
HIKEECIVPTPCYKPKEPLPVEAKLPWFKQAQELEEGAAV 520
SEEPS. 526
```

Signal peptide

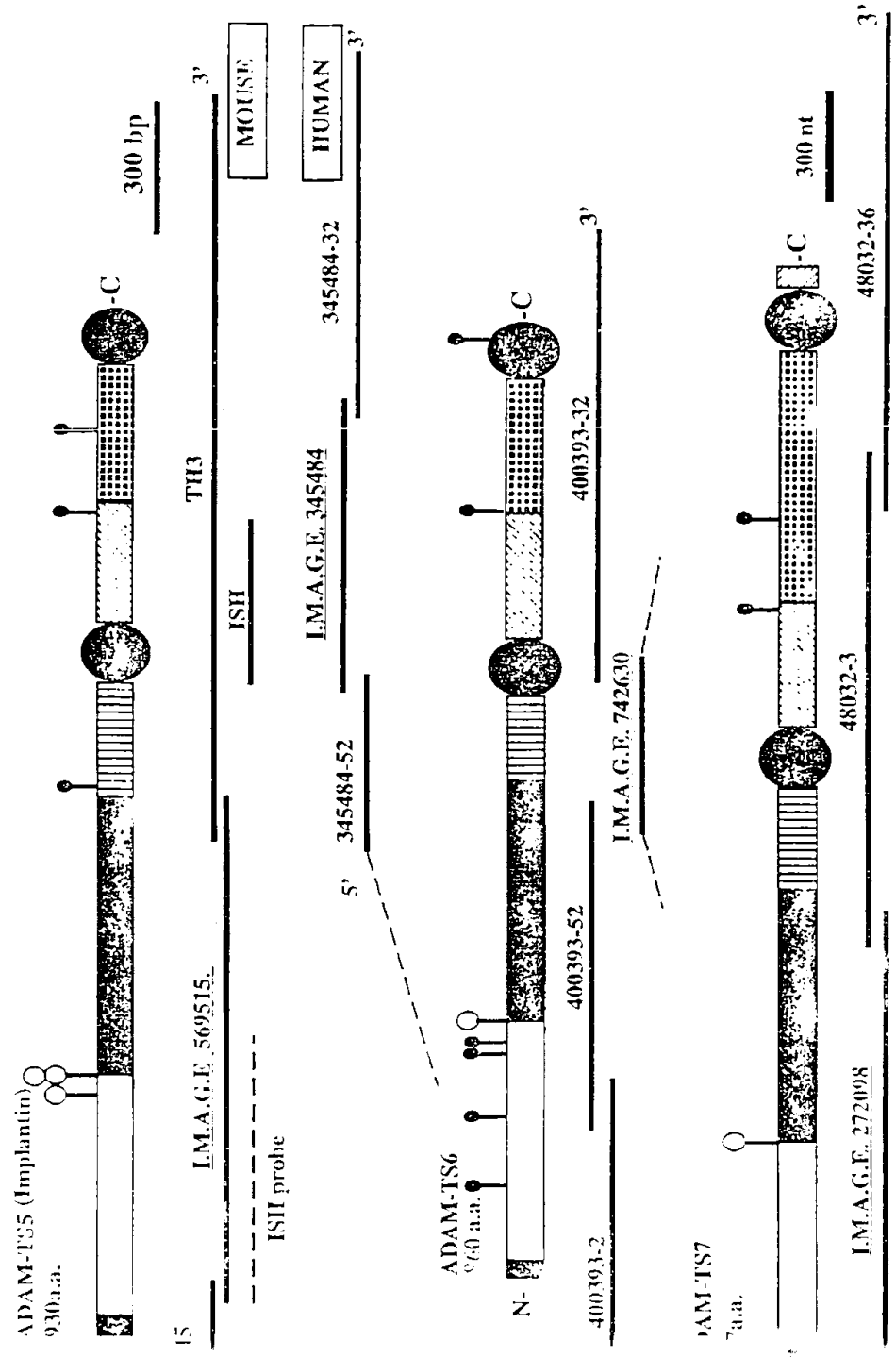
(C) YYPENIKPKPKLQE

(C) QELEEGAAV

C- terminal epitope for Ab

Similar to ADAM-TS family but lacks the prometalloprotease and disintegrin domain. Can [illegible] this may be a inhibitor of the

Fig. 12


ADAM-TS5 (Implantin)
930a.a.
-C
300 bp
3'
MOUSE
HUMAN
TH3
I.M.A.G.E. 569515
ISH
ISH probe
I.M.A.G.E. 345484
345484-32
5'
345484-52
ADAM-TS6
960 a.a.
N-
-C
400393-32
3'
400393-52
400393-2
I.M.A.G.E. 742630
ADAM-TS7
-C
300 nt
48032-3
48032-36
I.M.A.G.E. 272098
3'

a

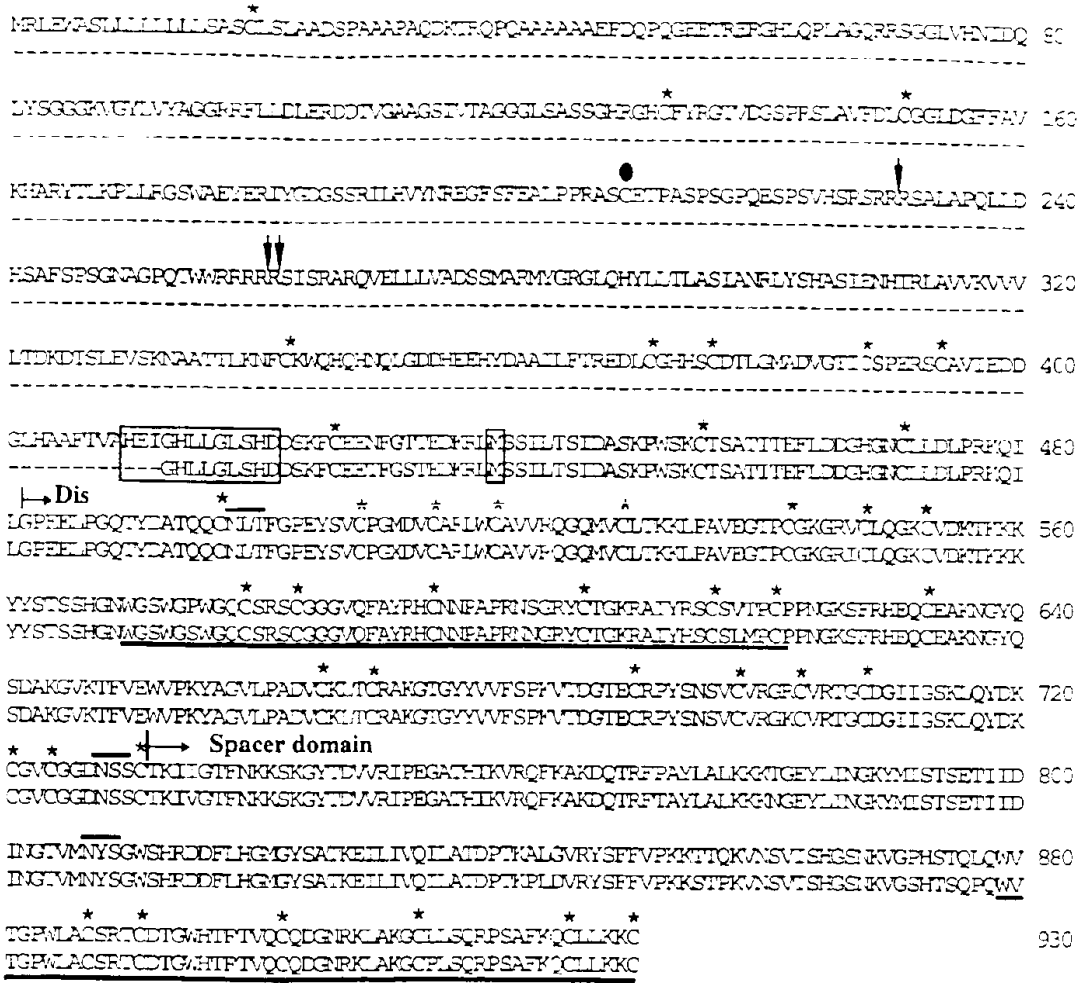
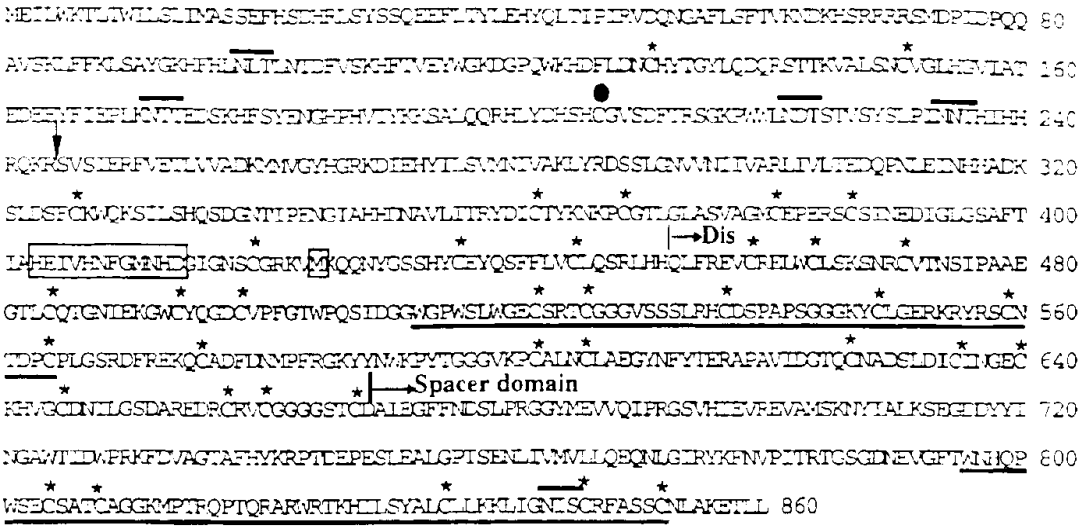


Fig. 13



C

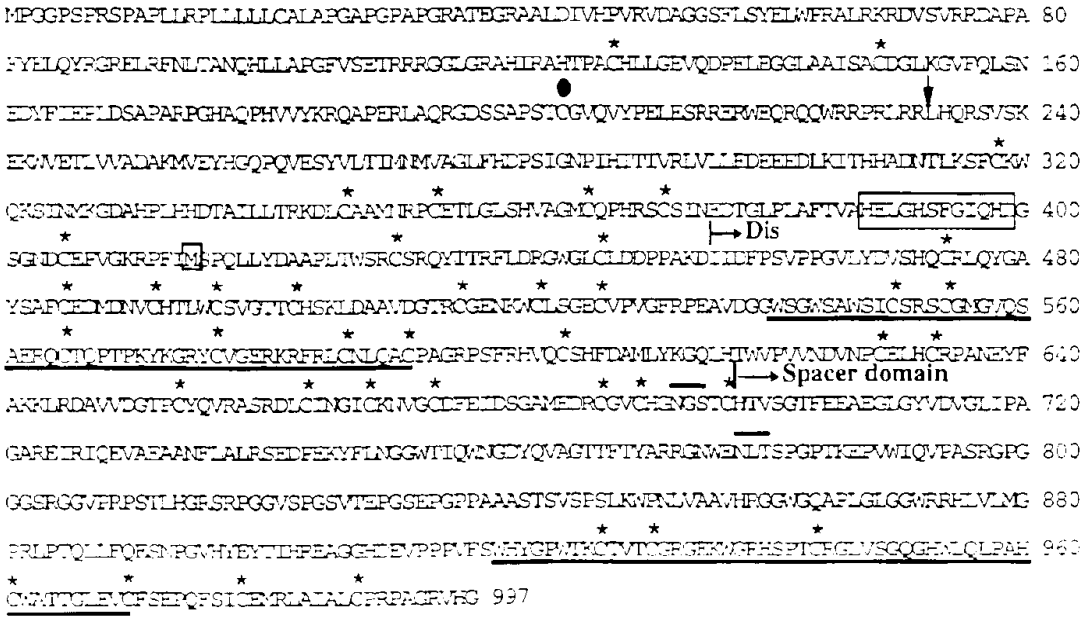


Fig. 13 (con't)

a

adamalysin II	HELGHNLGMEHD
atrolysin A	HELGHNLGMVHD
hADAM-9	HELGHNLGMNHD
hADAM-10	HEVGHNFGSPHD
hADAM-15	HELGHSLGLDHD
hADAM-17	HELGHNFGAEHD
mADAM-19	HEIGHNFGMSHD
mADAM-TS1	HELGHVFNMPHD
hADAM-TS2	HETGHVLGMEHD
hADAM-TS3	HETGHVLGMEHD
hADAM-TS4	HELGHVFNMLHD
mADAM-TS5	HEIGHLLGLSHD
hADAM-TS6	HEIVHNFGMNHD
hADAM-TS7	HELGHSFGIQHD

b

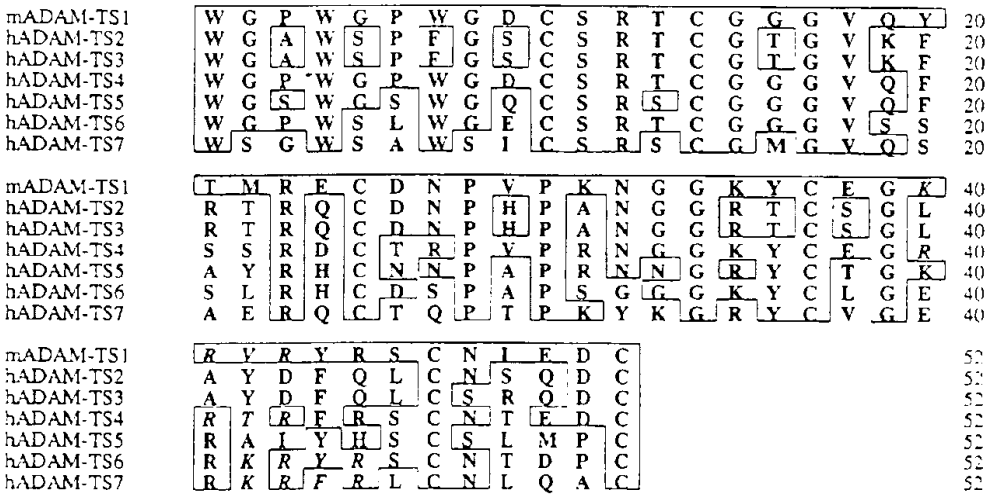


Fig. 13 (con't)

Fig. 14

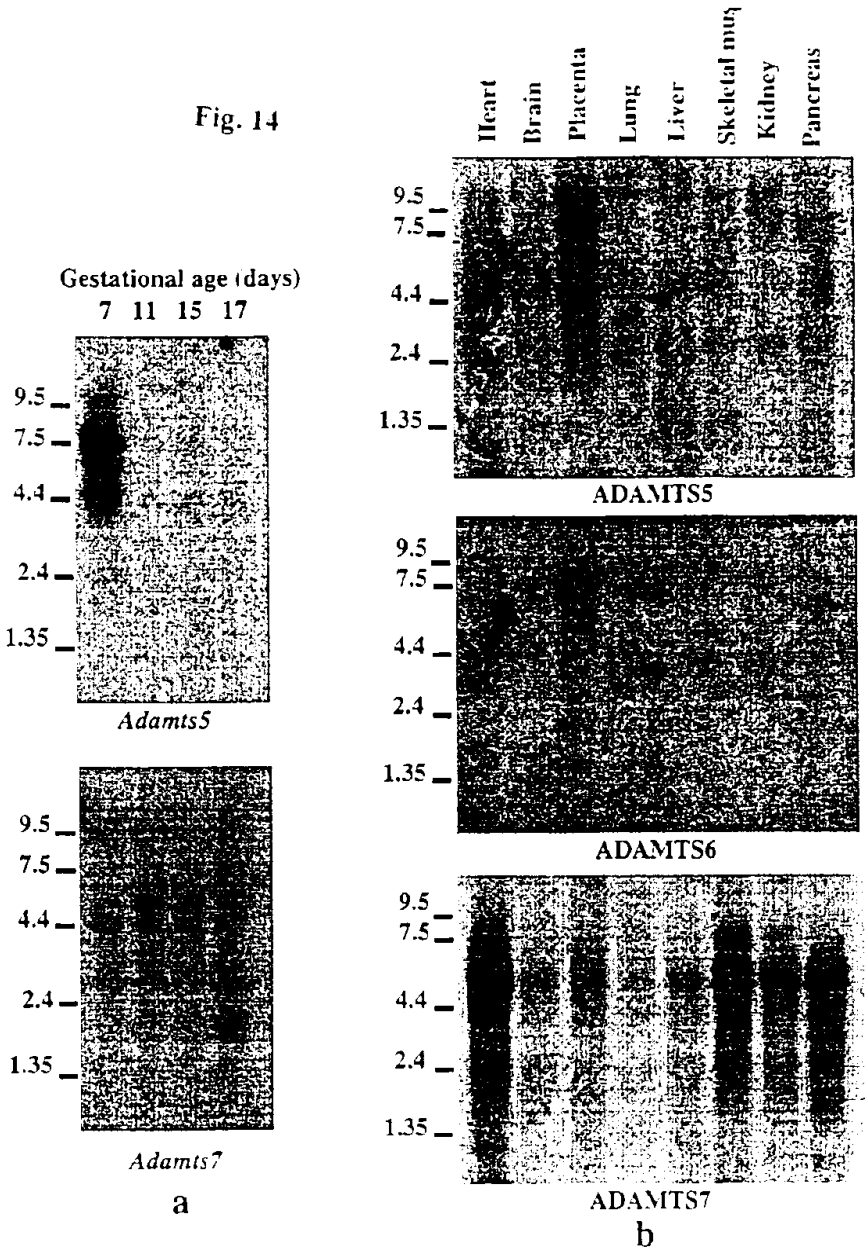

Gestational age (days)
7 11 15 17
9.5
7.5
4.4
2.4
1.35
Adamts5
Adamts7
a
Heart
Brain
Placenta
Lung
Liver
Skeletal muscle
Kidney
Pancreas
ADAMTS5
ADAMTS6
ADAMTS7
b

Fig. 15

ADAM-TS RELATED PROTEIN-1 (ADAM-TSR1)

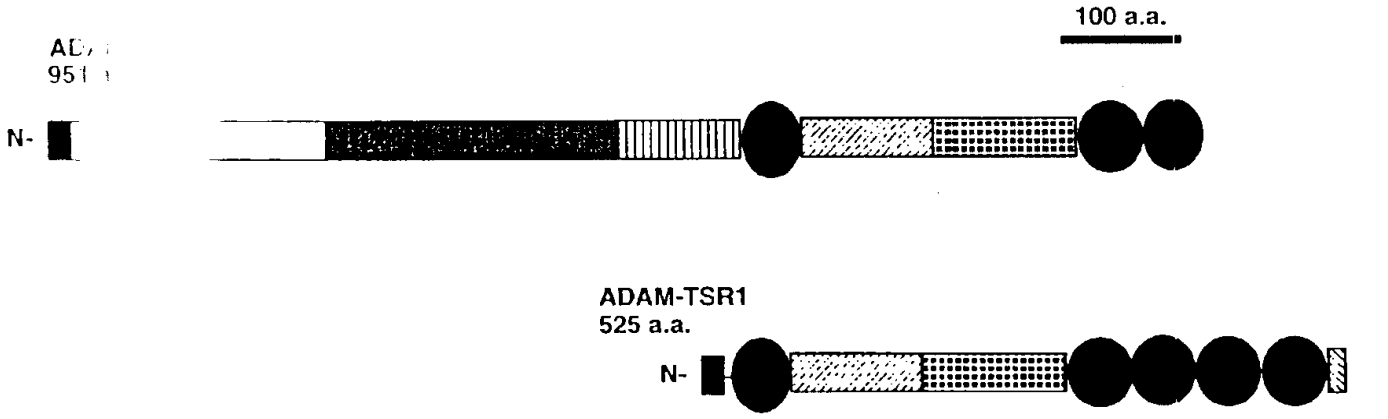


Fig. 15 (con't)

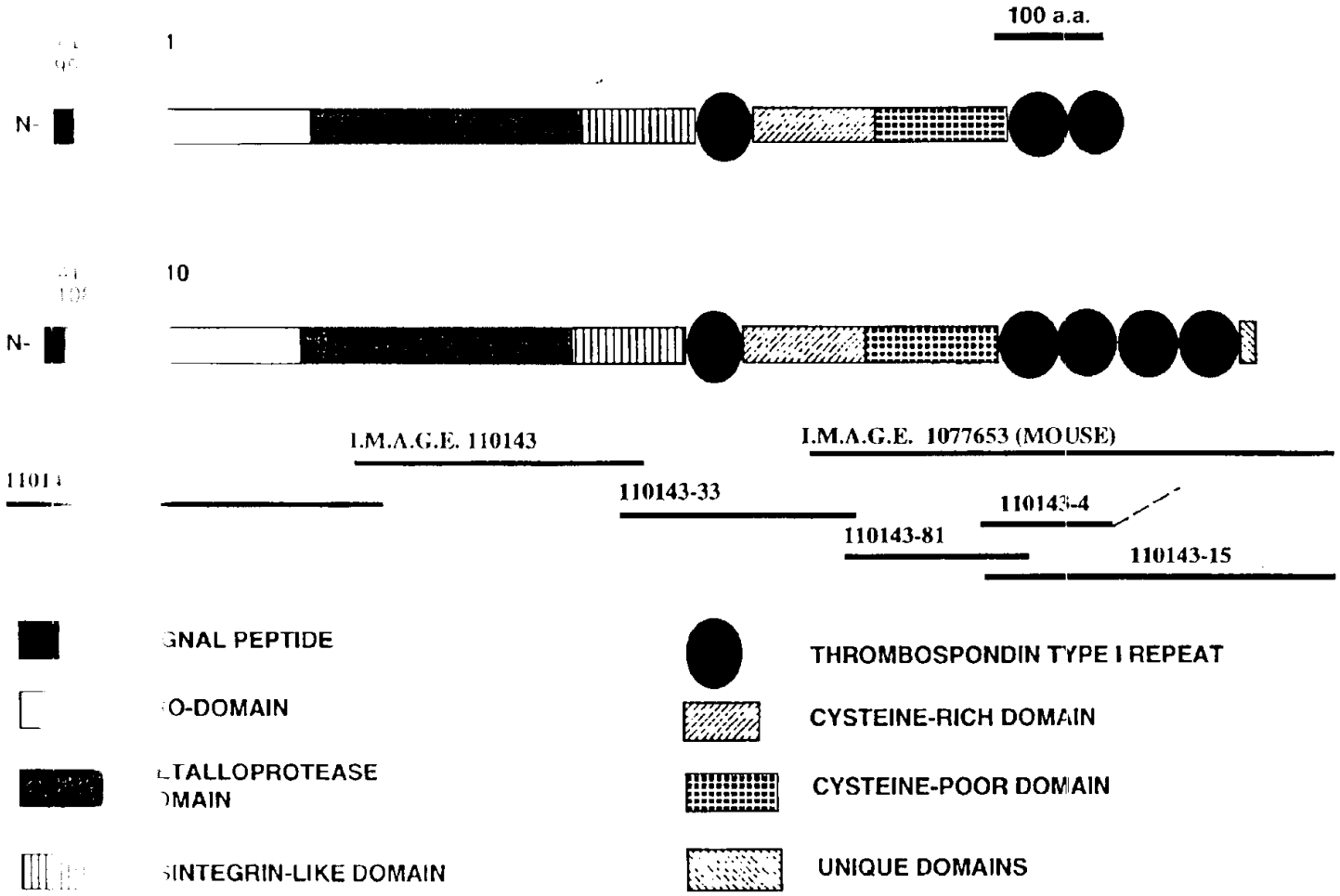

100 a.a.
1
N-
10
N-
I.M.A.G.E. 110143
I.M.A.G.E. 1077653 (MOUSE)
110143-33
110143-4
110143-81
110143-15
GNAL PEPTIDE
O-DOMAIN
TALLOPROTEASE
OMAIN
INTEGRIN-LIKE DOMAIN
THROMBOSPONDIN TYPE I REPEAT
CYSTEINE-RICH DOMAIN
CYSTEINE-POOR DOMAIN
UNIQUE DOMAINS

FIGURE 16

Pa

```
MSSCFVWPAMRSPSPPAWTTTGHCWPSRHLLP 40
GAAPRHGGHSRVPPLLQSGLASTHFLLNLTRSSRLLAGRV 80
SVEYWTREGLAWQRAARPHCLYAGHLQGQASSSHVAISTC 120
GGLHGLIVADEEEYLIEPLHGGPKGSRSPEESGPHVVYKR 160
SSLPHPHLDTACGVRDEKPWKGRPWWLRTLKPPPARPLGN 200
ETEFGQPGLKRSVSRERYVVETLVVADKMMVAYHGRRDVEQ 240
YVLAIMNIVAKLFQDSSLGSTVNILVTRLILLTEDQPTLE 280
ITHHAGHSLDSFCKWQKSIVNHSGHGNAIPENGVANHDTA 320
VLITRYDICIYKNKPCGTLGLARWAECVSAREAAASMRTL 360
AATSVHHCHEIGHTFGMNHDGVGNSCGARGQDPAKLMAAH 400
ITMTINPFVWSSCNRDYITSFLDSGLGLCLNNRPPRQDFV 440
YPTVAPGQAYDAEEQCRFQHGVKSRQCKYGEVCSELWCLS 480
KSNFCITNSIPAAEGTLCQTHTIDKGWCYKRVCVPFGSRP 520
EGVDGAWGPWTPWGDCSRTCGGGVSSSSRHCDSPRPTIGG 560
KYCLGERRRHRSCNTDDCPPGSQDFREVQCSEFDSIPFRG 600
KFYKWHTYRGGGVKACSLTSLAEGFNFYTERAAAVVDGTP 640
CRPDTVDICVSGECKHVGCDRVLGSDLREDKCRVCGGDGS 680
ACETIEGVFSPASPGAGYEDVVWIPKGSVHIFIQDLNLSL 720
SHLALPGDQESLLLEGLPGTPQPHRLPLAGTTFQLRQGPD 760
QVQSLEALGPLNASLIVMVLARTELPALRYRFNAPIARDS 800
LPPYSWHYAPWIKCSAQCAGGSQVQAVECRNQLDSSAVAP 840
HYCSAHSKLPKRQRACNTEPCPPDWVVGNWSLCSRSCDAG 880
VRSRSVVCQRRVSAAEEKALDDSACPQPRPPVLEACHGPT 920
CPPEWAALDWSECTPSCGPGLRHRVVLCKSADHRATLPPA 960
HCSPAAKPPATMRCNLRRCPPARWVAGEWGECSAQCGVGQ 1000
RQRSVRCTSHIGQASHECTEALRPPTTQQCEAKCDSPTPG 1040
DGPEECHDVNKVAYCPLVLKFQFCSRAYFRQMCCKTCQGH 1080
Created: Thursday, October 01, 1998 11:05 PM
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```
         10         20         30         40
         |          |          |          |
tcacgcacgccttccggtctcaagATGAGTTCCTGTCCAG 40
TCTGGAGAGCTTATGAGATCGCCTTCCCCACCCGCGTGGAC 80
CACAACGGGGCACTGCTGGCCTTCTCGCCACCTCCTCCCC 120
GGAGCAGCGCCGCGGCACGGGGGCCACAGCCGAGTCCCGC 160
CTCTTCTACAAAGTGGCCTCGCCAGCACCCACTTCCTGCT 200
```

FIGURE 16 (continued)

Pa

```
        210         220         230         240
GAACCTGACCCGCAGCTCCCGTCTACTGGCAGGGCGCGTC 240
TCCGTGGAGTACTGGACACGGGAGGGCCTGGCCTGGCAGA 280
GGGCGGCCCCGGCCCCACTGCCTCTACGCTGGTCACCTGCA 320
GGGCCAGGCCAGCAGCTCCCATGTGGCCATCAGCACCTGT 360
GGAGGCCTGCACGGCCTGATCGTGGCAGACGAGGAAGAGT 400
        410         420         430         440
ACCTGATTGAGCCCCTGCACGGTGGGCCCAAGGGTTCTCG 440
GAGCCCGGAGGAAAGTGGACCACATGTGGTGTACAAGCGT 480
TCCTCTCTGCGTCACCCCCACCTGGACACAGCCTGTGGAG 520
TGAGAGATGAGAAACCGTGGAAAGGGCGGCCATGGTGGCT 560
GCGGACCTTGAAGCCACCGCCTGCCAGACCCCTGGGGAAT 600
        610         620         630         640
GAAACAGAGCGTGGCCAGCCAGGCCTGAAGCGATCGGTCA 640
GCCGAGAGCGCTACGTGGAGACCCTGGTGGTGGCTGACAA 680
GATGATGGTGGCCTATCACGGGCGCCGGGATGTGGAGCAG 720
TATGTCCTGGCCATCATGAACATTGTTGCCAAACTTTTCC 760
AGGACTCGAGTCTGGGAAGCACCGTTAACATCCTCGTAAC 800
        810         820         830         840
TCGCCTCATCCTGCTCACGGAGGACCAGCCCACTCTGGAG 840
ATCACCCACCATGCCGGGAAGTCCCTAGACAGCTTCTGTA 880
AGTGGCAGAAATCCATCGTGAACCACAGCGGCCATGGCAA 920
TGCCATTCCAGAGAACGGTGTGGCTAACCATGACACAGCA 960
GTGCTCATCACACGCTATGACATCTGCATCTACAAGAACA 1000
        1010        1020        1030        1040
AACCCTGCGGCACACTAGGCCTGGCCCGGTGGCGGAATG 1040
TGTGACGCGAGAGAAGCTGCAGCGTCAATGAGGACATTG 1080
GCTGCCACAAGCGTTCACCATTGCCACGAGATCGGGCACA 1120
CATTCGGCATGAACCATGACGGCGTGGAAACAGCTGTGG 1160
```

FIGURE 16 (continued)

Pa

1210 1220 1230 1240

ATTACCATGAAGACCAACCCATTCGTGTGGTCATCCTGCA 1240
ACCGTGACTACATCACCAGCTTTCTAGACTCGGGCCTGGG 1280
GCTCTGCCTGAACAACCGGCCCCCCAGACAGGACTTTGTG 1320
TACCCGACAGTGGCACCGGGCCAAGCCTACGATGCAGATG 1360
AGCAATGCCGCTTTCAGCATGGAGTCAAATCGCGTCAGTG 1400

1410 1420 1430 1440

TAAATACGGGGAGGTCTGCAGCGAGCTGTGGTGTCTGAGC 1440
AAGAGCAACCGGTGCATCACCAACAGCATCCCGCCGCCCG 1480
AGGGCACGCTGTGCCAGACGCACACCATCGACAAGGGGTG 1520
GTGCTACAAACGGGTCTGTGTCCCCTTTGGGTCGCGCCCA 1560
GAGGGTGTGGACGGAGCCTGGGGGCCGTGGACTCCATGGG 1600

1610 1620 1630 1640

GCGACTGCAGCCGGACCTGTGGCGGCGGCGTGTCCTCTTC 1640
TAGTCGTCACTGCGACAGCCCCAGGCCAACCATCGGGGGC 1680
AAGTACTGTCTGGGTGAGAGAAGGCGGCACCGCTCCTGCA 1720
ACACGGATGACTGTCCCCCTGGCTCCCAGGACTTCAGAGA 1760
AGTGCAGTGTTCTGAATTTGACAGCATCCCTTTCCGTGGG 1800

1810 1820 1830 1840

AAATTCTACAAGTGGAAAACGTACCGGGGAGGGGGCGTGA 1840
AGGCCTGCTCGCTCACGAGCCTAGCGGAAGGCTTCAACTT 1880
CTACACGGAGACGGCGGCAGCCGTGGTGGACGGGACACCC 1920
TGCCGTCCAGACACGGTGGACATTTGCGTCAGTGGCGAAT 1960
GCAAGCACGTGGGCTGCGACCGAGTCCTGGGCTCCGACCT 2000

2010 2020 2030 2040

GCGGAGGACAAGTGCCGAGTGTGTGGCGGTGACGGCAGT 2040
GCTTGCGAGACCATCGAGGGCGTCTTCAGCCCAGCCTCAC 2080
CTGGGGCCGGTACGAGGATGTCGTCTGGATTCCCAAAGG 2120
CTCCGTCCACATCTTCATCCAGGATCTGAACCTCTCTCTC 2160

FIGURE 16 (continued)

Pa

```
        2210        2220        2230        2240
TGGAGGGGCTGCCTGGGACCCCCCAGCCCCACCGTCTGCC 2240
TCTAGCTGGGACCACCTTTCAACTGCGACAGGGGCCAGAC 2280
CAGGTCCAGAGCCTCGAAGCCCTGGGACCGATTAATGCAT 2320
CTCTCATCGTCATGGTGCTGGCCCGGACCGAGCTGCCTGC 2360
CCTCCGCTACCGCTTCAATGCCCCCATCGCCCGTGACTCG 2400
        2410        2420        2430        2440
CTGCCCCCCTACTCCTGGCACTATGCGCCCTGGACCAAGT 2440
GCTCGGCCCAGTGTGCAGGCGGTAGCCAGGTGCAGGCGGT 2480
GGAGTGCCGCAACCAGCTGGACAGCTCCGCGGTCGCCCCC 2520
CACTACTGCAGTGCCCACAGCAAGCTGCCCAAAAGGCAGC 2560
GCGCCTGCAACACGGAGCCTTGCCCTCCAGACTGGGTTGT 2600
        2610        2620        2630        2640
AGGGAACTGGTCGCTCTGCAGCCGCAGCTGCGATGCAGGC 2640
GTGCGCAGTCGCTCGGTCGTGTGCCAGCGCCGCGTCTCTG 2680
CCGCGGAGGAGAAGGCGCTGGACGACAGCGCATGCCCGCA 2720
GCCGCGCCCACCTGTACTGGAGGCCTGCCACGGCCCCACT 2760
TGCCCTCCGGAGTGGGCGGCCCTCGACTGGTCTGAGTGCA 2800
        2810        2820        2830        2840
CCCCCAGCTGCGGGCCGGGCCTCCGCCACCGCGTGGTCCT 2840
TTGCAAGAGCGCAGACCACCGCGCCACGCTGCCCCGGCG 2880
CACTGCTCACCCGCCGCCAAGCCACCGGCCACCATGCGCT 2920
GCAACTTGCGCCGCTGCCCCCCGGCCCGCTGGGTGGCTGG 2960
CGAGTGGGGTGAGTGCTCTGCACAGTGCGGCGTCGGGCAG 3000
        3010        3020        3030        3040
CGGCAGCGCTCGGTGCGCTGCACCAGCCACACGGGCCAGG 3040
CGTCGCACGAGTGCACGGAGGCCCTGCGGCCGCCCACCAC 3080
GCAGCAGTGTGAGGCCAAGTGCGACAGCCCAACCCCCGGG 3120
GACGGCCCTGAAGAGTGCAAGGATGTGAACAAGGTCGCCT 3160
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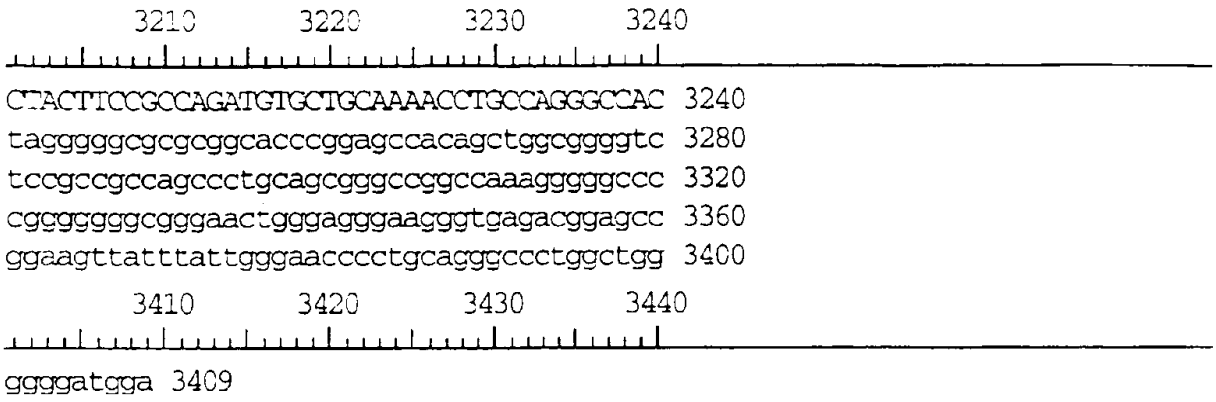
FIGURE 16 (continued)



Pa

FIGURE 17

Molecular Weight 216301.30 Daltons
1934 Amino Acids
234 Strongly Basic(+) Amino Acids (K,R)
216 Strongly Acidic(-) Amino Acids (D,E)
477 Hydrophobic Amino Acids (A,I,L,F,W,V)
657 Polar Amino Acids (N,C,Q,S,T,Y)

7.734 Isolectric Point
24.102 Charge at PH 7.0

MQFVSWATLLTLLVRDLAEMGSPDAAAAVRKDRLHPRQVKLLETLSEYEIVSPIRVNALG 60
EPFPTNVHFKRTRRSINSATDPWPAFASSSSSSTSPQAHYRLSAFGQQFLFNLTANAGFI 120
APLFTVTLLGTPGVNQTKFYSEEEAELKHCFYKGYVNINSEHTAVISLCSGMLGTFRSHD 180
GGYFIEPLQSMDEQEDEEEQNKPHIIYRRSAPQREPSTGRHACDTSEHKNRHSKDKKKTR 240
ARKWGERINLAGDVAALNSGLATEAFSAYGNKTDNTREKRTHRRTKRFLSYPRFVEVLVV 300
ADNRMVSYHGENLQHYILTLMSIVASIYKDPSIGNLINIVIVNLIVIHNEQDGPSISFNA 360
QTTLKNFCQWQHSNSPGGIHHDTAVLLTRQDICRAHDKCDITLGLAELGTICDPYRSCSIS 420
EDSGLSTAFTIAHELGHVFNMPHDDNNKCKEEGVKSPQHVMAPTLNFYTNPWMWSKCSRK 480
YITEFLDTGYGECLLNEPESRPYPLPVQLPGILYNVNKQCELIFGPGSQVCPYMMQCRRL 540
WCNVVNGVHKGCRTQHTPWADGTECEPGKHCKYGFCVPKEMDVPVTDGSWGSWSPFGTCS 600
RTCGGGIKTAIRECNRPEPKNGGKYCVGRRMKFKSCNTEPCLKQKRDFRDEQCAHFDGKH 660
FNTNGLLPNVFWVPKYSGILMKDRCKLFCRVAGNTAYYQLRDRVIDGTPCGQDTNDICVQ 720
GLCRQAGCDHVLNSKARRDKCGVCGGDNSSCKTVAGTFNTVHYGYNTVVRIPAGATNIDV 780
RQHSFSGEIDDDNYLALSSSKGEFLLNGNFVVTMAKREIRIGNAVVEYSGSETAVERINS 840
TDRIEQELLLQVLSVGKLYNPDVRYSFNIPIEDKPQQFYWNSHGPWQACSKPCQGERKRK 900
LVCTRESDQLTVSDQRCDRLPQPGHITEPCGTGCDLRWHVASRSECSAQCGLGYRTLDIY 960
CAKYSRLDGKTEKVDDGFCSSHPKPSNREKCSGECNTGGWRYSAWTECSKSCDGGTQRRR 1020
AICVNTRNDVLDDSKCTHQEKVTIQRCSEFPCPQWKSGDWSECLVTCGKGHKHRQVWCQF 1080
GEDFLNDRWCDPETKPTSMQTCQQPECASWQAGPWVQCSVTCGQGYQLRAVKCIIGTYMS 1140
VVDENDCNAATRPTDTQDCELPSCHPPPAAPETRRSTYSAPRTQWRFGSWTPCSATCGKG 1200
TRMRYVSCFDENGSVADESACATLPRPVAKEECSVTPCGQWKALDWSSCSVTCGQGRATR 1260
QVMCVNYSDHVTDRSECDQDYIPETDQDCSMSPCPQRTPDSGLACHPFQNEDYRPRSASP 1320
SRTHVLGGNQWRTGPWGACSSTCAGGSQRRVVVCQDENGYTANDCVERIKPDEQPACESG 1380
PCPQWAYGNWGECTKLCGGGIRTRLVVCQRSNGERFPDLSCEILDKPPDREQCNTHACPH 1440
DAAWSTGPWSSCSVSCGRGHKQRNVYCMAKDGSHLESDYCKHLAKPHGHRKCRGGRCPKW 1500
KAGAWSQCSVSCGRGVQQRHVGCQIGTHKIAFETECNPYTRPESECECQGPRCPLYIWRA 1560
EEVQECTNTCGEGSRYRKVVCVDINKNEVHGARCDVSKRPVDRESCSLQPCEYVWITGEW 1620

[illegible]

FIGURE 17 (continued)

```
DCYSAAKCPQGRFSINLYGTGLSLTESARWLSQGNYAVSDIKKSPDGTRVVGKCGGYCGK 1920
CTPSSGTGLEVRVL 1934

          10         20         30         40
          |          |          |          |
tggggcagcggagggaggggtgggaagcaccATGCAGTT 40
TGTATCCTGGGCCACACTGCTAACGCTCCTGGTGCGGGAC 80
CTGGCCGAGATGGGGAGCCCAGACGCCGCGGCGGCCGTGC 120
GCAAGGACAGGCTGCACCCGAGGCAAGTGAAATTATTAGA 160
GACCCTGAGCGAATACGAAATCGTGTCTCCCATCCGAGTG 200
         210        220        230        240
         |          |          |          |
AACGCTCTCGGAGAACCCTTTCCCACGAACGTCCACTTCA 240
AAAGAACGCGACGGAGCATTAACTCTGCCACTGACCCCTG 280
GCCTGCCTTCGCCTCCTCCTCTTCCTCCTCTACCTCCCCC 320
CAGGCGCATTACCGCCTCTCTGCCTTCGGCCAGCAGTTTC 360
TATTTAATCTCACCGCCAATGCCGGATTTATCGCTCCACT 400
         410        420        430        440
         |          |          |          |
GTTCACTGTCACCCTCCTCGGGACGCCCGGGGTGAATCAG 440
ACCAAGTTTTATTCCGAAGAGGAAGCGGAACTCAAGCACT 480
GTTTCTACAAAGGCTATGTCAATACCAACTCCGAGCACAC 520
GGCCGTCATCAGCCTCTGCTCAGGAATGCTGGGCACATTC 560
CGGTCTCATGATGGGGGTTATTTTATTGAACCACTACAGT 600
         610        620        630        640
         |          |          |          |
CTATGGATGAACAAGAAGATGAAGAGGAACAAAACAAACC 640
CCACATCATTTATAGGCGCAGCGCCCCCCAGAGAGAGCCC 680
TCAACAGGAAGGCATGCATGTGACACCTCAGAACACAAAA 720
ATAGGCACAGTAAAGACAAGAAGAAAACCAGAGCAAGAAA 760
ATGGGGAGAAAGGATTAACCTGGCTGGTGACGTAGCAGCA 800
         810        820        830        840
         |          |          |          |
[illegible]
GAGAAAACCTTCAACACCAGATTTTAACTTTAATGTCAAT 1000
```

FIGURE 17 (continued)

Pa

1010 1020 1030 1040

TGTAGCCTCTATCTATAAAGACCCAAGTATTGGAAATTTA 1040
ATTAATATTGTTATTGTGAACTTAATTGTGATTCATAATG 1080
AACAGGATGGGCCTTCCATATCTTTTAATGCTCAGACAAC 1120
ATTAAAAAACTTTTGCCAGTGGCAGCATTCGAACAGTCCA 1160
GGTGGAATCCATCATGATACTGCTGTTCTCTTAACAAGAC 1200

1210 1220 1230 1240

AGGATATCTGCAGAGCTCACGACAAATGTGATACCTTAGG 1240
CCTGGCTGAACTGGGAACCATTTGTGATCCCTATAGAAGC 1280
TGTTCTATTAGTGAAGATAGTGGATTGAGTACAGCTTTTA 1320
CGATCGCCCATGAGCTGGGCCATGTGTTTAACATGCCTCA 1360
TGATGACAACAACAAATGTAAAGAAGAAGGAGTTAAGAGT 1400

1410 1420 1430 1440

CCCCAGCATGTCATGGCTCCAACACTGAACTTCTACACCA 1440
ACCCCTGGATGTGGTCAAAGTGTAGTCGAAAATATATCAC 1480
TGAGTTTTTAGACACTGGTTATGGCGAGTGTTTGCTTAAC 1520
GAACCTGAATCCAGACCCTACCCTTTGCCTGTCCAACTGC 1560
CAGGCATCCTTTACAACGTGAATAAACAATGTGAATTGAT 1600

1610 1620 1630 1640

TTTTGGACCAGGTTCTCAGGTGTGCCCATATATGATGCAG 1640
TGCAGACGGCTCTGGTGCAATAACGTCAATGGAGTACACA 1680
AAGGCTGCCGGACTCAGCACACACCCTGGGCCGATGGGAC 1720
GGAGTGCGAGCCTGGAAAGCACTGCAAGTATGGATTTTGT 1760
GTTCCCAAAGAAATGGATGTCCCCGTGACAGATGGATCCT 1800

1810 1820 1830 1840

GGGGAAGTTGGAGTCCCTTTGGAACCTGCTCCAGAACATG 1840
TGGAGGGGGCATCAAAACAGCCATTCGAGAGTGCAACAGA 1880
CCAGAACCAAAAAATGGTGCAAAATACTGTGTAGGACGTA 1920
GAATGAAACTTAAGTCCTGCAACACGGAGCCATGTCTCAA 1960

FIGURE 17 (continued)

```
        2010        2020        2030        2040
GACGGGAAGCATTTTAACATCAACGGTCTGCTTCCCAATG 2040
TGCGCTGGGTCCCTAAATACAGTGGAATTCTGATGAAGGA 2080
CCGGTGCAAGTTGTTCTGCAGAGTGGCAGGGAACACAGCC 2120
TACTATCAGCTTCGAGACAGAGTGATAGATGGAACTCCTT 2160
GTGGCCAGGACACAAATGATATCTGTGTCCAGGGCCTTTG 2200
        2210        2220        2230        2240
CCGGCAAGCTGGATGCGATCATGTTTTAAACTCAAAAGCC 2240
CGGAGAGATAAATGCGGGGTTTGTGGTGGCGATAATTCTT 2280
CATGCAAAACAGTGGCAGGAACATTTAATACAGTACATTA 2320
TGGTTACAATACTGTGGTCCGAATTCCAGCTGGTGCTACC 2360
AATATTGATGTGCGGCAGCACAGTTTCTCAGGGGAAACAG 2400
        2410        2420        2430        2440
ACGATGACAACTACTTAGCTTTATCAAGCAGTAAAGGTGA 2440
ATTCTTGCTAAATGGAAACTTTGTTGTCACAATGGCCAAA 2480
AGGGAAATTCGCATTGGGAATGCTGTGGTAGAGTACAGTG 2520
GGTCCGAGACTGCCGTAGAAAGAATTAACTCAACAGATCG 2560
CATTGAGCAAGAACTTTTGCTTCAGGTTTTGTCGGTGGGA 2600
        2610        2620        2630        2640
AAGTTGTACAACCCCGATGTACGCTATTCTTTCAATATTC 2640
CAATTGAAGATAAACCTCAGCAGTTTTACTGGAACAGTCA 2680
TGGGCCATGGCAAGCATGCAGTAAACCCTGCCAAGGGGAA 2720
CGGAAACGAAAACTTGTTTGCACCAGGGAATCTGATCAGC 2760
TTACTGTTTCTGATCAAAGATGCGATCGGCTGCCCCAGCC 2800
        2810        2820        2830        2840
TGGACACATTACTGAACCCTGTGGTACAGGCTGTGACCTG 2840
AGGTGGCATGTTGCCAGCAGGAGTGAATGTAGTGCCCAGT 2880
GTGGCTTGGGTTACCGCACATTGGACATCTACTGTGCCAA 2920
ATATAGCAGGCTGGATGGGAAGACTGAGAAGGTTGATGAT 2960
```

FIGURE 17 (continued)

Pa

3010 3020 3030 3040

AATGCTCAGGGGAATGTAACACGGGTGGCTGGCGCTATTC 3040
TGCCTGGACTGAATGTTCAAAAAGCTGTGACGGTGGGACC 3080
CAGAGGAGAAGGGCTATTTGTGTCAATACCCGAAATGATG 3120
TACTGGATGACAGCAAATGCACACATCAAGAGAAAGTTAC 3160
CATTCAGAGGTGCAGTGAGTTCCCTTGTCCACAGTGGAAA 3200

3210 3220 3230 3240

TCTGGAGACTGGTCAGAGTGCTTGGTCACCTGTGGAAAAG 3240
GGCATAAGCACCGCCAGGTCTGGTGTCAGTTTGGTGAAGA 3280
TCGATTAAATGATAGAATGTGTGACCCTGAGACCAAGCCA 3320
ACATCTATGCAGACTTGTCAGCAGCCGGAATGTGCATCCT 3360
GGCAGGCGGGTCCCTGGGTACAGTGCAGTGTCACTTGTGG 3400

3410 3420 3430 3440

ACAGGGATACCAGCTAAGAGCAGTGAAATGCATCATTGGG 3440
ACTTATATGTCAGTGGTAGATGACAATGACTGTAATGCAG 3480
CAACTAGACCAACTGATACCCAGGACTGTGAATTACCATC 3520
ATGTCATCCTCCCCCAGCTGCCCCCGGAAACGAGGAGAAGC 3560
ACATACAGTGCACCAAGAACCCAGTGGCGATTTGGGTCTT 3600

3610 3620 3630 3640

GGACCCCATGCTCAGCCACTTGTGGGAAAGGTACCCGGAT 3640
GAGATACGTCAGCTGCCGAGATGAGAATGGCTCTGTGGCT 3680
GACGAGAGTGCCTGTGCTACCCTGCCTAGACCAGTGGCAA 3720
AGGAAGAATGTTCTGTGACACCCTGTGGGCAATGGAAGGC 3760
CTTGGACTGGAGCTCTTGCTCTGTGACCTGTGGGCAAGGT 3800

3810 3820 3830 3840

AGGGCAACCCGGCAAGTGATGTGTGTCAACTACAGTGACC 3840
ACGTGATCGATCGGAGTGAGTGTGACCAGGATTATATCGC 3880
AGAAACTGACCAGGACTGTTCCATGTCACCATGCCCTCAA 3920
AGGACCCCAGACAGTGGCTTAGCTCAGCACCCTTCCAAA 3960

FIGURE 17 (continued)

4010 4020 4030 4040

CCATGTGCTCGGTGGAAACCAGTGGAGAACTGGCCCCTGG 4040
GGAGCATGTTCCAGTACCTGTGCTGGCGGATCCCAGCGGC 4080
GTGTTGTTGTATGTCAGGATGAAAATGGATACACCGCAAA 4120
CGACTGTGTGGAGAGAATAAAACCTGATGAGCAAAGAGCC 4160
TGTGAATCCGGCCCTTGTCCTCAGTGGGCTTATGGCAACT 4200

4210 4220 4230 4240

GGGGAGAGTGCACTAAGCTGTGTGGTGGAGGCATAAGAAC 4240
AAGACTGGTGGTCTGTCAGCGGTCCAACGGTGAACGGTTT 4280
CCAGATTTGAGCTGTGAAATTCTTGATAAACCTCCCGATC 4320
GTGAGCAGTGTAACACACATGCTTGTCCACACGACGCTGC 4360
ATGGAGTACTGGCCCTTGGAGCTCGTGTTCTGTCTCTTGT 4400

4410 4420 4430 4440

GGTCGAGGGCATAAACAACGAAATGTTTACTGCATGGCAA 4440
AAGATGGAAGCCATTTAGAAAGTGATTACTGTAAGCACCT 4480
GGCTAAGCCACATGGGCACAGAAAGTGCCGAGGAGGAAGA 4520
TGCCCCAAATGGAAAGCTGGCGCTTGGAGTCAGTGCTCTG 4560
TGTCCTGTGGCCGAGGCGTACAGCAGAGGCATGTGGGCTG 4600

4610 4620 4630 4640

TCAGATCGGAACACACAAAATAGCCAGAGAGACCGAGTGC 4640
AACCCATACACCAGACCGGAGTCGGAATGCGAATGCCAAG 4680
GCCCACGGTGTCCCCTTTACACTTGGAGGGCAGAGGAATG 4720
GCAAGAATGCACCAAGACCTGCGGCGAAGGCTCCAGGTAC 4760
CGCAAGGTGGTGTGTGTGGATGACAACAAAAACGAGGTGC 4800

4810 4820 4830 4840

ATGGGGCACGCTGTGACGTGAGCAAGCGGCCGGTGGACCG 4840
TGAAAGCTGTAGTTTGCAACCCTGCGAGTATGTCTGGATC 4880
ACAGGAGAATGGTCAGAGTGCTCAGTGACCTGTGGAAAAG 4920
GCTACAAACAAAGGCTTGTCTCGTGCAGCGAGATTTACAC 4960

FIGURE 17 (continued)

Pa

5010 5020 5030 5040

AACTGCCCAGGCACGCAGCCCCCCAGTGTTCACCCCTGTT 5040
ACCTGAGGGAGTGCCCTGTCTCGGCCACCTGGAGAGTTGG 5080
CAACTGGGGGAGCTGCTCAGTGTCTTGTGGTGTTGGAGTG 5120
ATGCAGAGATCTGTGCAATGTTTAACCAATGAGGACCAAC 5160
CCAGCCACTTATGCCACACTGATCTGAAGCCAGAAGAACG 5200

5210 5220 5230 5240

AAAAACCTGCCGTAATGTCTATAACTGTGAGTTACCCCAG 5240
AATTGCAAGGAGGTAAAAAGACTTAAAGGTGCCAGTGAAG 5280
ATGGTGAATATTTCCTGATGATTAGAGGAAAGCTTCTGAA 5320
GATATTCTGTGCGGGGATGCACTCTGACCACCCCAAAGAG 5360
TACGTGACACTGGTGCATGGAGACTCTGAGAATTTCTCCG 5400

5410 5420 5430 5440

AGGTTTATGGGCACAGGTTACACAACCCAACAGAATGTCC 5440
CTATAACGGGAGCCGGTGCGATGACTGCCAATGTCGGAAG 5480
GATTACACGGCCGCTGGGTTTTCCAGTTTTCAGAAAATCA 5520
GAATAGACCTGACCAGCATGCAGATAATCACCACTGACTT 5560
ACAGTTTGCAAGGACAAGCGAAGGACATCCCGTCCCTTTT 5600

5610 5620 5630 5640

GCTACAGCCGGGGATTGCTACAGCGCTGCCAAGTGCCCAC 5640
AGGGTCGTTTTAGCATCAACCTTTATGGAACCGGCTTGTC 5680
TTTAACTGAATCTGCCAGATGGATATCACAAGGGAATTAT 5720
GCTGTCTCTGACATCAAGAAGTCCGCCGGATGGTACCCGAG 5760
TCGTAGGGAAATGCGGTGGTTACTGTGGAAAATGCACTCC 5800

5810 5820 5830 5840

ATCCTCTGGTACTGGCTGGAGGTGCGAGTTTTAtagcta 5840
aggtgctttgaagaggaagccattatggatggatgaagga 5880
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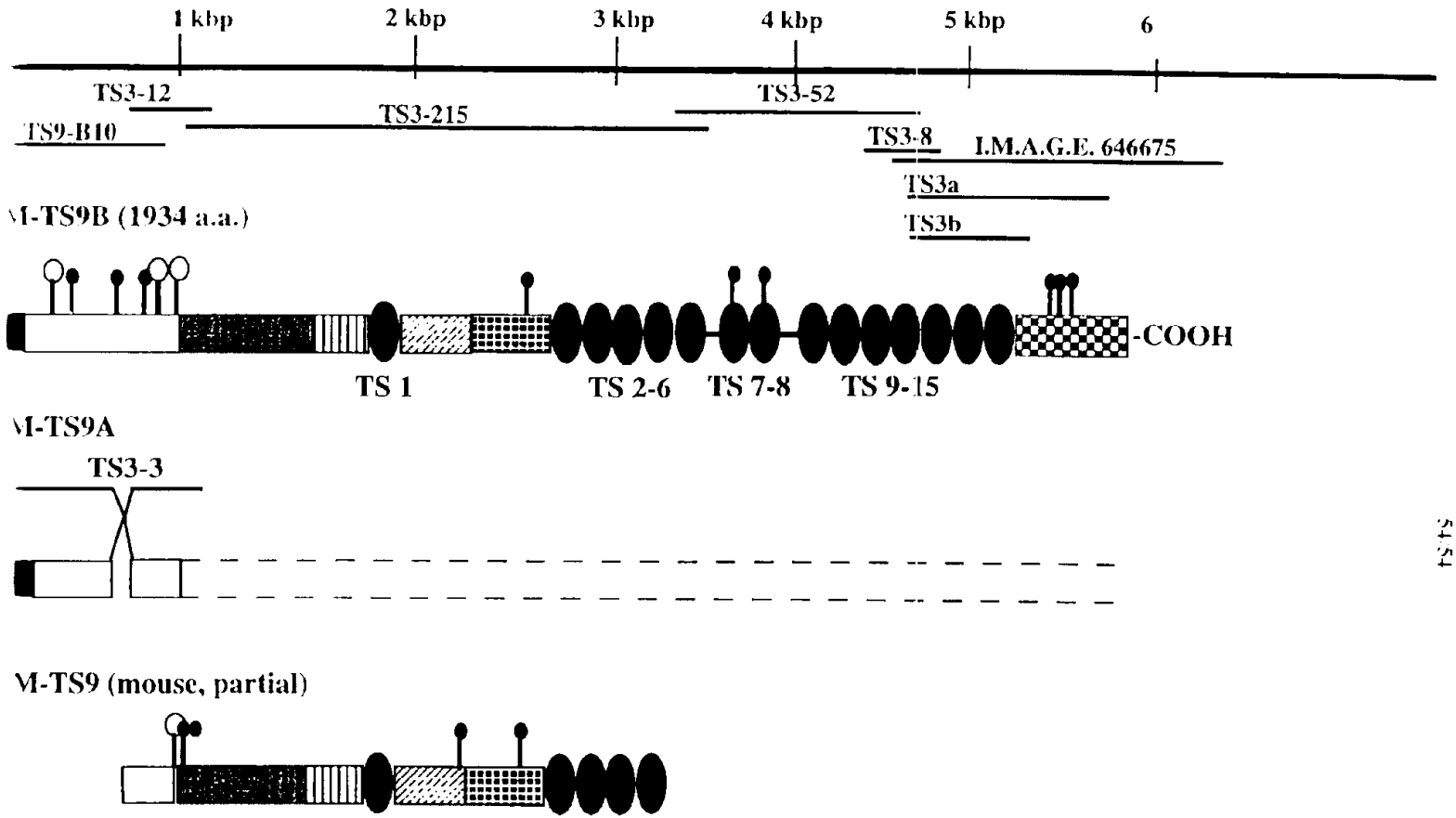
atgtgtgtgtgtgtttgtgtgtgacttgtatgcttgtgtg 5960

FIG



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Leu Leu Leu Leu Ser Ala Ser Cys Leu Ser Leu Ala Ala Asp Ser Pro
15 20 25

gcc gcg gca cct gcc cag gat aaa acc agg cag cct cag gct gca gca 146
Ala Ala Ala Pro Ala Gln Asp Lys Thr Arg Gln Pro Gln Ala Ala Ala
30 35 40

gcg gcc gcc gag ccg gac cag ccg cag ggg gag gaa aca cgg gag cga 194
Ala Ala Ala Glu Pro Asp Gln Pro Gln Gly Glu Glu Thr Arg Glu Arg
45 50 55

ggc cat tta caa ccc ttg gcc ggg cag cgc agg agc ggc ggg ctg gtc 242
Gly His Leu Gln Pro Leu Ala Gly Gln Arg Arg Ser Gly Gly Leu Val
60 65 70 75

cat aat ata gac caa ctc tac tct ggc ggt ggc aaa gtg ggc tat ctt 290
His Asn Ile Asp Gln Leu Tyr Ser Gly Gly Gly Lys Val Gly Tyr Leu
80 85 90

gtc tat gcg ggc ggc cgg agg ttc ctg ctg gac ctg gag aga gat gac 338
Val Tyr Ala Gly Gly Arg Arg Phe Leu Leu Asp Leu Glu Arg Asp Asp
95 100 105

aca gtg ggt gct gct ggt agc atc gtt act gta gga ggg ggg ctg agc 386
Thr Val Gly Ala Ala Gly Ser Ile Val Thr Ala Gly Gly Gly Leu Ser
110 115 120

[illegible]
Gly Ser Ile Arg Asn Leu Ala Val Ile Asp Ile Gly Glu Gly Ile
140 145 150 155

```
   ggc ttc ttt gca gtc aag cat gcg cgc tac act cta aag cca ctc ctg   530
   Gly Phe Phe Ala Val Lys His Ala Arg Tyr Thr Leu Lys Pro Leu Leu
                   160                 165                 170
5
   cgt ggg tcc tgg gca gag tat gaa cga att tat ggg gat gga tct tcc   578
   Arg Gly Ser Trp Ala Glu Tyr Glu Arg Ile Tyr Gly Asp Gly Ser Ser
               175                 180                 185

10 cgc atc ctg cat gtc tac aac cgc gag ggc ttt agc ttc gag gcc ctg   626
   Arg Ile Leu His Val Tyr Asn Arg Glu Gly Phe Ser Phe Glu Ala Leu
           190                 195                 200

   ccg cca cgc gcc agt tgc gag act cct gca tcc cca tct ggg ccc caa   674
15 Pro Pro Arg Ala Ser Cys Glu Thr Pro Ala Ser Pro Ser Gly Pro Gln
       205                 210                 215

   gag agc ccc tcg gtg cac agt aga tct agg aga cgc tca gcg ctg gcc   722
   Glu Ser Pro Ser Val His Ser Arg Ser Arg Arg Arg Ser Ala Leu Ala
20 220                 225                 230                 235

   ccg cag ctg ctg gac cac tca gct ttc tcg cca tct ggg aac gcg gga   770
   Pro Gln Leu Leu Asp His Ser Ala Phe Ser Pro Ser Gly Asn Ala Gly
                   240                 245                 250
25
   cct cag act tgg tgg agg cgt agg cgc cgt tcc atc tcc agg gcc cgc   818
   Pro Gln Thr Trp Trp Arg Arg Arg Arg Arg Ser Ile Ser Arg Ala Arg
               255                 260                 265

30 cag gtg gag ctc ctc ttg gtg gct gac tcg tcc atg gcc agg atg tat   866
   Gln Val Glu Leu Leu Leu Val Ala Asp Ser Ser Met Ala Arg Met Tyr
           270                 275                 280

   ggg cgg ggc ctg cag cat tac ctg ctg acc atg gcc tcc atc gcc aac   914
35 Gly Arg Gly Leu Gln His Tyr Leu Leu Thr Met Ala Ser Ile Ala Asn
       285                 290                 295

   agg ctg tac agt cat gca agc att gag aac cac atc cgc ctg gcg gtg   962
   Arg Leu Tyr Ser His Ala Ser Ile Glu Asn His Ile Arg Leu Ala Val
40 300                 305                 310                 315

   gtg aag gtg gtg gtg ctg acg gac aag gac acg agt ctg gag gtg agc  1010
   Val Lys Val Val Val Leu Thr Asp Lys Asp Thr Ser Leu Glu Val Ser
                   320                 325                 330
45
   aag aat ggg gcc acg acc ctc aag aac ttt tgc aaa tgg cag cac cag  1058
   Lys Asn Ala Ala Thr Thr Leu Lys Asn Phe Cys Lys Trp Gln His Gln
               335                 340                 345

50 cat aac cag cta ggg gat gat cac gaa gag cac tat gat gca gcc atc  1106
   His Asn Gln Leu Gly Asp Asp His Glu Glu His Tyr Asp Ala Ala Ile
           350                 355                 360

   ctg ttc acc cga gag gat tta tgt ggg cat cat tca tgt gac acc ctg  1154
55 Leu Phe Thr Arg Glu Asp Leu Cys Gly His His Ser Cys Asp Thr Leu
       365                 370                 375

   agc aac agt aac ttt ggg acc ata tct aag gac gtt aaa ttg gcc agc tat gga  1202

   att ggg tat cta ctt gga att tct gat gcc gtt cca ttt atg tgt gaa
```

```
   Ile Gly His Leu Leu Gly Leu Ser His Asp Asp Ser Lys Phe Cys Glu
                415                 420                 425

   gag aac ttc ggt act aca gaa gac aag cgt tta atg tct tca atc ctt   1346
 5 Glu Asn Phe Gly Thr Thr Glu Asp Lys Arg Leu Met Ser Ser Ile Leu
            430                 435                 440

   acc agc atc gat gca tcc aag ccc tgg tcc aaa tgc acg tca gcc acc   1394
   Thr Ser Ile Asp Ala Ser Lys Pro Trp Ser Lys Cys Thr Ser Ala Thr
10      445                 450                 455

   atc aca gaa ttc ctg gat gat ggt cat ggt aat tgt ttg cta gac cta   1442
   Ile Thr Glu Phe Leu Asp Asp Gly His Gly Asn Cys Leu Leu Asp Leu
   460                 465                 470                 475
15
   cca cgg aag cag att ttg ggt ccc gag gaa ctc cca gga cag acc tac   1490
   Pro Arg Lys Gln Ile Leu Gly Pro Glu Glu Leu Pro Gly Gln Thr Tyr
                480                 485                 490

20 gat gcc acc cag cag tgc aac ttg aca ttt ggg cct gag tac tcg gtg   1538
   Asp Ala Thr Gln Gln Cys Asn Leu Thr Phe Gly Pro Glu Tyr Ser Val
                495                 500                 505

   tgc cct ggc atg gat gtc tgt gcg cgg ctg tgg tgt gct gtg gtg cgc   1586
25 Cys Pro Gly Met Asp Val Cys Ala Arg Leu Trp Cys Ala Val Val Arg
            510                 515                 520

   caa ggc caa atg gtg tgt ctg acc aag aag ctg ccg gct gtg gag ggc   1634
   Gln Gly Gln Met Val Cys Leu Thr Lys Lys Leu Pro Ala Val Glu Gly
30      525                 530                 535

   act ccc tgt ggg aag gga aga gtc tgc ctt caa ggc aaa tgt gtg gac   1682
   Thr Pro Cys Gly Lys Gly Arg Val Cys Leu Gln Gly Lys Cys Val Asp
   540                 545                 550                 555
35
   aaa act aag aaa aaa tat tac tcg aca tca agc cat gga aat tgg ggg   1730
   Lys Thr Lys Lys Lys Tyr Tyr Ser Thr Ser Ser His Gly Asn Trp Gly
                560                 565                 570

40 tcc tgg ggc ccc tgg ggt cag tgt tct cgc tca tgc ggg gga gga gtg   1778
   Ser Trp Gly Pro Trp Gly Gln Cys Ser Arg Ser Cys Gly Gly Gly Val
            575                 580                 585

   cag ttt gcc tac cgc cat tgt aat aac cct gca cct cga aac agt ggc   1826
45 Gln Phe Ala Tyr Arg His Cys Asn Asn Pro Ala Pro Arg Asn Ser Gly
        590                 595                 600

   cgc tac tgc acc ggg aag agg gcc ata tac cgt tcc tgc agt gtt acc   1874
   Arg Tyr Cys Thr Gly Lys Arg Ala Ile Tyr Arg Ser Cys Ser Val Thr
50      605                 610                 615

   ccc tgc cca ccc aat ggt aaa tct ttt cgc cat gag cag tgt gaa gcc   1922
   Pro Cys Pro Pro Asn Gly Lys Ser Phe Arg His Glu Gln Cys Glu Ala
   620                 625                 630                 635
55
   aaa aat ggc tat cag tct gat gca aaa gga gtc aaa aca ttt gta gaa   1970
   Lys Asn Gly Tyr Gln Ser Asp Ala Lys Gly Val Lys Thr Phe Val Glu
                640                 645                 650

   ttt tgc gat gtg act gtg gat ctg gag ttc gtc aag gac ttt gtg gtg   2018
   Phe Cys Asp Val Thr Val Asp Leu Glu Phe Val Lys Asp Phe Val Val
            655                 660                 665

   gag agc atc tac cct ggc tac ctg gcc atg gag gcc acc ggg gag ctg   2066
   Glu Ser Ile Tyr Pro Gly Tyr Leu Ala Met Glu Ala Thr Gly Glu Leu
        670                 675                 680

   ctg gcc aag aag aag gtg gac ctg gcc gtg gtg gtg gtg gac gag gcc   2114
   Leu Ala Lys Lys Lys Val Asp Leu Ala Val Val Val Val Asp Glu Ala
   685                 690                 695

   aac gcc acc tac gcc ggc tcc gtg cag tcc gtg cac gac ctg ggc ctg   2162
   Asn Ala Thr Tyr Ala Gly Ser Val Gln Ser Val His Asp Leu Gly Leu
   700                 705                 710                 715

   ctg ggc gag gtg cgc aac ctg agc ttc tcc ggc gtg cgc aac ttc ggc   2210
   Leu Gly Glu Val Arg Asn Leu Ser Phe Ser Gly Val Arg Asn Phe Gly
                720                 725                 730

   tcc atc gag gag gtg gtg gac ctg gag ccc gac gtg ccc gtg cgc acc   2258
   Ser Ile Glu Glu Val Val Asp Leu Glu Pro Asp Val Pro Val Arg Thr
            735                 740                 745

   ttc gtg cag aac gcc aac ccc ctg cgc gag gtg gac gcc tcc gtg ctg   2306
   Phe Val Gln Asn Ala Asn Pro Leu Arg Glu Val Asp Ala Ser Val Leu
        750                 755                 760

   cac ccc gag cgc gtg ggc ggc ttc ctg gtg ctg gac ctg ctg cgc ccc   2354
   His Pro Glu Arg Val Gly Gly Phe Leu Val Leu Asp Leu Leu Arg Pro
   765                 770                 775

   ctg gtg ctg ccc gac ggc tcc ctg ctg gac ctg gag gac gtg gac gcc   2402
   Leu Val Leu Pro Asp Gly Ser Leu Leu Asp Leu Glu Asp Val Asp Ala
   780                 785                 790                 795

   ctg ctg gag ctg ggc tgg ctg cgc gag gtg gcc ggc gtg tgg ccc ccc   2450
   Leu Leu Glu Leu Gly Trp Leu Arg Glu Val Ala Gly Val Trp Pro Pro
                800                 805                 810

   gcc ggc ggc ctg ctg gag gtg gcc ctg ccc gag tac gtg gag gcc ctg   2498
   Ala Gly Gly Leu Leu Glu Val Ala Leu Pro Glu Tyr Val Glu Ala Leu
            815                 820                 825

   ggc aac ctg cac tac gcc cgc ggc cag acc ccc gtg gcc aag gcc atg   2546
   Gly Asn Leu His Tyr Ala Arg Gly Gln Thr Pro Val Ala Lys Ala Met
        830                 835                 840

   atg cag ctg ttc tcc gac gag atc ctg ccc ctg gtg cgc gcc tac gcc   2594
   Met Gln Leu Phe Ser Asp Glu Ile Leu Pro Leu Val Arg Ala Tyr Ala
   845                 850                 855

   ggc ggc ggc ccc gag ctg gag tgg gtg gac ggc aag ccc ctg ctg cgc   2642
   Gly Gly Gly Pro Glu Leu Glu Trp Val Asp Gly Lys Pro Leu Leu Arg
   860                 865                 870                 875

   ctg cgc ggc ggc gag ctg ggc aag cgc gtg ctg gac gtg gag gtg ccc   2690
   Leu Arg Gly Gly Glu Leu Gly Lys Arg Val Leu Asp Val Glu Val Pro
                880                 885                 890

   ctg gcc ctg ctg ctg gtg ctg ggc ggc gag ggc ctg ctg tgg ctg cgc   2738
   Leu Ala Leu Leu Leu Val Leu Gly Gly Glu Gly Leu Leu Trp Leu Arg
            895                 900                 905

   ctg ctg ctg gac ctg gcc ctg ctg ccc ctg ggc ctg ctg ctg ctg ctg   2786
   Leu Leu Leu Asp Leu Ala Leu Leu Pro Leu Gly Leu Leu Leu Leu Leu
        910                 915                 920

   ctg gcc ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   2834
   Leu Ala Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
   925                 930                 935

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   2882
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
   940                 945                 950                 955

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   2930
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
                960                 965                 970

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   2978
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
            975                 980                 985

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3026
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
        990                 995                1000

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3074
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1005                1010                1015

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3122
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1020                1025                1030                1035

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3170
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               1040                1045                1050

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3218
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
           1055                1060                1065

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3266
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
       1070                1075                1080

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3314
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1085                1090                1095

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3362
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1100                1105                1110                1115

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3410
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               1120                1125                1130

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3458
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
           1135                1140                1145

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3506
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
       1150                1155                1160

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3554
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1165                1170                1175

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3602
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1180                1185                1190                1195

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3650
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               1200                1205                1210

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3698
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
           1215                1220                1225

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3746
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
       1230                1235                1240

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3794
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1245                1250                1255

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3842
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1260                1265                1270                1275

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3890
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               1280                1285                1290

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3938
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
           1295                1300                1305

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   3986
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
       1310                1315                1320

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4034
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1325                1330                1335

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4082
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1340                1345                1350                1355

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4130
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               1360                1365                1370

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4178
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
           1375                1380                1385

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4226
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
       1390                1395                1400

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4274
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1405                1410                1415

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4322
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1420                1425                1430                1435

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4370
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               1440                1445                1450

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4418
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
           1455                1460                1465

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4466
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
       1470                1475                1480

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4514
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1485                1490                1495

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4562
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1565                1570                1575

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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
  1580                1585                1590                1595

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   4850
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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  1725                1730                1735

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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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       1790                1795                1800

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           1935                1940                1945

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  2205                2210                2215

   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg   6722
   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
               2240                2245                2250

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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu Leu
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   ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg ctg
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10 cta caa tat gac aag tgt gga gtg tgc gga ggg gat aac tcc agt tgt   2210
   Leu Gln Tyr Asp Lys Cys Gly Val Cys Gly Gly Asp Asn Ser Ser Cys
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   aca aag att atc gga acc ttc aat aaa aaa agc aag ggt tat act gac   2258
15 Thr Lys Ile Ile Gly Thr Phe Asn Lys Lys Ser Lys Gly Tyr Thr Asp
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   Lys Ala Lys Asp Gln Thr Arg Phe Pro Ala Tyr Leu Ala Leu Lys Lys
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   Lys Thr Gly Glu Tyr Leu Ile Asn Gly Lys Tyr Met Ile Ser Thr Ser
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30 gag acc atc atc gac atc aat ggt acc gtc atg aac tac agt gga tgg   2450
   Glu Thr Ile Ile Asp Ile Asn Gly Thr Val Met Asn Tyr Ser Gly Trp
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   agc cac aga gat gat ttt tta cat ggg atg ggc tat tca gcc aca aaa   2498
35 Ser His Arg Asp Asp Phe Leu His Gly Met Gly Tyr Ser Ala Thr Lys
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   gaa atc ctg atc gtg cag atc ctt gcc aca gac cca act aaa gcg cta   2546
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40         830                 835                 840

   ggc gtc cgt tac agc ttt ttt gtt ccc aag aag acc act caa aaa gta   2594
   Gly Val Arg Tyr Ser Phe Phe Val Pro Lys Lys Thr Thr Gln Lys Val
       845                 850                 855
45
   aac tct gtc atc agc cat ggc agt aac aag gtg gga cca cac tct aca   2642
   Asn Ser Val Ile Ser His Gly Ser Asn Lys Val Gly Pro His Ser Thr
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50 cag ttg cag tgg gtg aca ggt cca tgg ctg gcc tgc tct agg acc tgt   2690
   Gln Leu Gln Trp Val Thr Gly Pro Trp Leu Ala Cys Ser Arg Thr Cys
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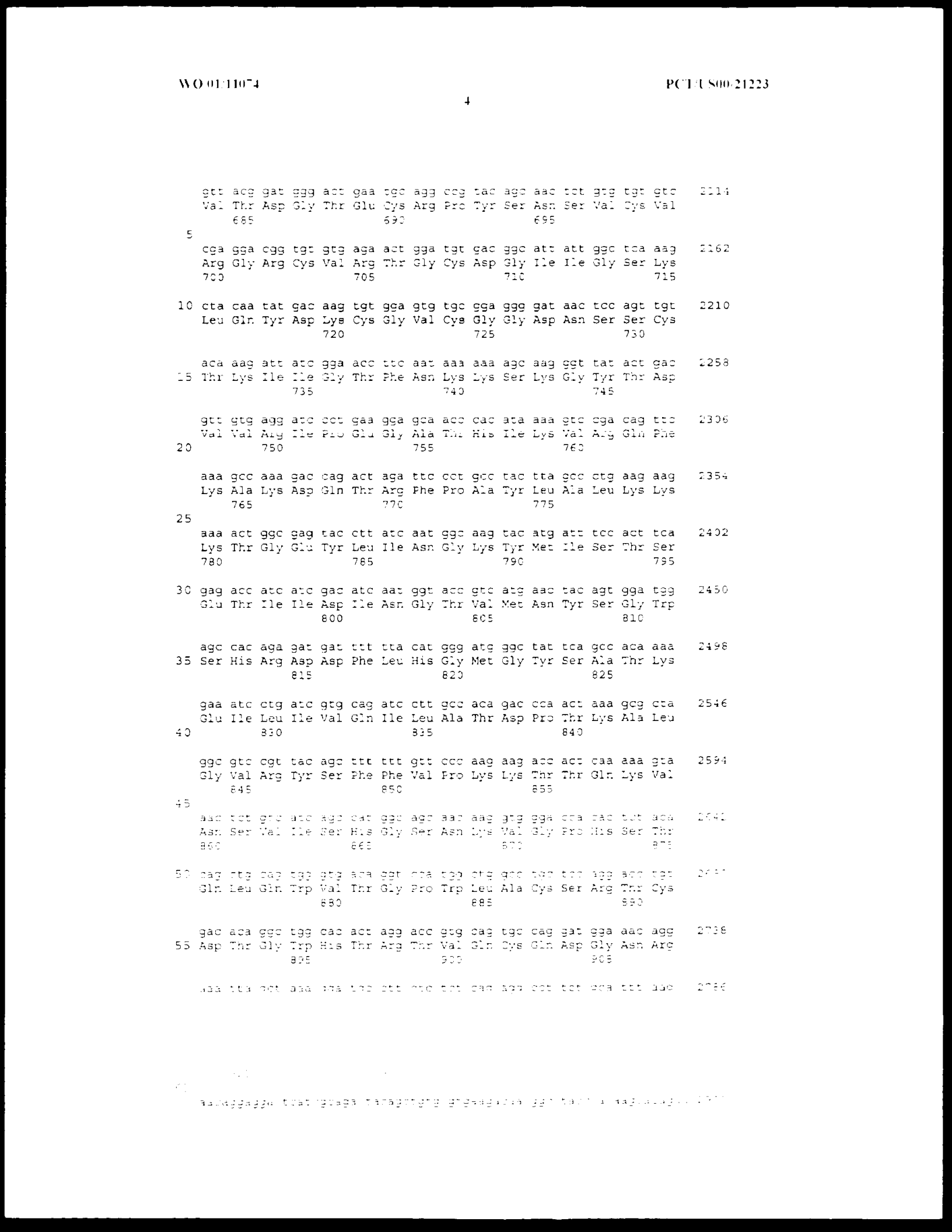

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Gln	Asp	Lys	Thr	Arg	Gln	Pro	Gln	Ala	Ala	Ala	Ala	Ala	Ala	Glu	Pro
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Arg	Arg	Phe	Leu	Leu	Asp	Leu	Glu	Arg	Asp	Asp	Thr	Val	Gly	Ala	Ala
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Gly	Ser	Ile	Val	Thr	Ala	Gly	Gly	Gly	Leu	Ser	Ala	Ser	Ser	Gly	His
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Lys	His	Ala	Arg	Tyr	Thr	Leu	Lys	Pro	Leu	Leu	Arg	Gly	Ser	Trp	Ala
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Glu	Tyr	Glu	Arg	Ile	Tyr	Gly	Asp	Gly	Ser	Ser	Arg	Ile	Leu	His	Val
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Tyr	Asn	Arg	Glu	Gly	Phe	Ser	Phe	Glu	Ala	Leu	Pro	Pro	Arg	Ala	Ser
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Glu Gly Ala Thr His Ile Lys Val Arg Gln Phe Lys Ala Lys Asp Gln
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   gcc tat ggc aag cac ttt cat cta aac ttg act ctc aac aca gat ttt   339
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   gtg tcc aaa cat ttt aca gta gaa tat tgg ggg aaa gat gga ccc cag   387
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   tgg aaa cat gat ttt tta gac aac tgt cat tac aca gga tat ttg caa   435
   Trp Lys His Asp Phe Leu Asp Asn Cys His Tyr Thr Gly Tyr Leu Gln
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   Asp Gln Arg Ser Thr Thr Lys Val Ala Leu Ser Asn Cys Val Gly Leu
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   Pro His Val Ile Tyr Lys Lys Ser Ala Leu Gln Gln Arg His Leu Tyr
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   gat cac tct cat tgt ggg gtt tcg gat ttc aca aga agt ggc aaa cct   675
   Asp His Ser His Cys Gly Val Ser Asp Phe Thr Arg Ser Gly Lys Pro
           205                 210                 215

40 tgg tgg ctg aat gac act cca ctg ttt ctt att cac tac caa att aac   723
   Trp Trp Leu Asn Asp Thr Pro Leu Phe Leu Ile His Tyr Gln Ile Asn
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   aac aca cat atc cac cac aga cag aag aga tca gtg agc att gaa cgg   771
45 Asn Thr His Ile His His Arg Gln Lys Arg Ser Val Ser Ile Glu Arg
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   Phe Val Glu Thr Leu Val Val Ala Asp Lys Met Met Val Gly Tyr His
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   ggc cgc aaa gac att gaa cat tac att ttg agt gtg atg aat att gtt   867
   Gly Arg Lys Asp Ile Glu His Tyr Ile Leu Ser Val Met Asn Ile Val
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   gcc aaa ctt tac cgt gat tcc agc cta gga aac gtt gtg aat att ata   915
   Ala Lys Leu Tyr Arg Asp Ser Ser Leu Gly Asn Val Val Asn Ile Ile
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   att gcc cac cac gat aat gca gtt ctt att act aga tat gat atc tgc   1107
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10 act tat aaa aat aag ccc tgt gga aca ctg ggc ttg gcc tct gtg gct   1155
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   gga atg tgt gag cct gaa agg agc tgc agc att aat gaa gac att ggc   1203
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   Leu Gly Ser Ala Phe Thr Ile Ala His Glu Ile Val His Asn Phe Gly
20 395                 400                 405                 410

   atg aac cat gat gga att gga aat tct tgt gga cga aag gtc atg aag   1299
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   Gln Gln Asn Tyr Gly Ser Ser His Tyr Cys Glu Tyr Gln Ser Phe Phe
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   aaa ggg tgg tgt tat cag gga gat tgt gtt cct ttt ggc act tgg ccc   1539
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50 agt agg acc tgt ggg gga ggc gtt tcc tca tcc cta aga cat tgt gac   1635
   Ser Arg Thr Cys Gly Gly Gly Val Ser Ser Ser Leu Arg His Cys Asp
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   Pro Ala Val Ile Asp Gly Thr Gln Cys Asn Ala Asp Ser Leu Asp Ile
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   tgc att aat gga gaa tgc aag cac gta ggc tgt gat aat att ttg gga  1971
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   tct gat gct agg gaa gat aga tgt cga gtc tgt gga ggg ggc gga agc  2019
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20 aca tgt gat gcc att gaa ggg ttc ttc aat gat tca ctg ccc agg gga  2067
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   ggc tac atg gaa gtg gtg cag ata cca aga ggc tct gtt cac att gaa  2115
25 Gly Tyr Met Glu Val Val Gln Ile Pro Arg Gly Ser Val His Ile Glu
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   gga gat gat tac tat att aat ggt gcc tgg act att gac tgg cct agg  2211
   Gly Asp Asp Tyr Tyr Ile Asn Gly Ala Trp Thr Ile Asp Trp Pro Arg
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          765                 770                 775

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   Asn Val Pro Ile Thr Arg Thr Gly Ser Gly Asp Asn Glu Val Gly Phe
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Leu
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Lys His Ser Arg Arg Arg Arg Ser Met Asp Pro Ile Asp Pro Gln Gln
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Ser Lys His Phe Ser Tyr Glu Asn Gly His Pro His Val Ile Tyr Lys
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Val Ser Asp Phe Thr Arg Ser Gly Lys Pro Trp Trp Leu Asn Asp Thr
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Leu Trp Pro Arg Ala Leu Arg Lys Arg Asp Val Ser Val Arg Arg Asp
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   2510                2515                2520                2525

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   2750                2755                2760                2765

635 gac gac gac gac gac gac gac gac gac gac gac gac gac gac gac gac  8355
   Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp Asp
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   Asp
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   aca ctc tgg tgc tct gtg ggg acc acc tgt cac tcc aag ctg gat gca   1539
   Thr Leu Trp Cys Ser Val Gly Thr Thr Cys His Ser Lys Leu Asp Ala
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5
   gct gtg gac ggc acc cgg tgt ggg gag aat aag tgg tgt ctc agt ggg   1587
   Ala Val Asp Gly Thr Arg Cys Gly Glu Asn Lys Trp Cys Leu Ser Gly
   510                 515                 520                 525

10 gag tgc gta ccc gtg ggc ttc cgg ccc gag gcc gtg gat ggt ggc tgg   1635
   Glu Cys Val Pro Val Gly Phe Arg Pro Glu Ala Val Asp Gly Gly Trp
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   tct ggc tgg agc gcc tgg tcc atc tgc tca cgg agc tgt ggc atg ggc   1683
15 Ser Gly Trp Ser Ala Trp Ser Ile Cys Ser Arg Ser Cys Gly Met Gly
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   gta cag agc gcc gag cgg cag tgc acg cag cct acg ccc aaa tac aaa   1731
   Val Gln Ser Ala Glu Arg Gln Cys Thr Gln Pro Thr Pro Lys Tyr Lys
20         560                 565                 570

   ggc aga tac tgt gtg ggt gag cgc aag cgc ttc cgc ctc tgc aac ctg   1779
   Gly Arg Tyr Cys Val Gly Glu Arg Lys Arg Phe Arg Leu Cys Asn Leu
       575                 580                 585
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   cag gcc tgc cct gct ggc cgc ccc tcc ttc cgc cac gtc cag tgc agc   1827
   Gln Ala Cys Pro Ala Gly Arg Pro Ser Phe Arg His Val Gln Cys Ser
   590                 595                 600                 605

30 cac ttt gac gct atg ctc tac aag ggc cag ctg cac aca tgg gtg ccc   1875
   His Phe Asp Ala Met Leu Tyr Lys Gly Gln Leu His Thr Trp Val Pro
                   610                 615                 620

   gtg gtc aat gac gtg aac ccc tgc gag ctg cac tgc cgg ccc gcg aat   1923
35 Val Val Asn Asp Val Asn Pro Cys Glu Leu His Cys Arg Pro Ala Asn
               625                 630                 635

   gag tac ttt gcc aag aag ctg cgg gac gcc tgt gtc gat ggc acc ccc   1971
   Glu Tyr Phe Ala Lys Lys Leu Arg Asp Ala Cys Val Asp Gly Thr Pro
40         640                 645                 650

   tgc tac cag gtc cga gcc agc cgg gac ctc tgc atc aac ggc atc tgt   2019
   Cys Tyr Gln Val Arg Ala Ser Arg Asp Leu Cys Ile Asn Gly Ile Cys
       655                 660                 665
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   aag aac gtg ggc tgt gac ttc gag att gac tcc ggt gct atg gag gac   2067
   Lys Asn Val Gly Cys Asp Phe Glu Ile Asp Ser Gly Ala Met Glu Asp
   670                 675                 680                 685

50 cgc tgt cgt gtg tgc cac ggc aac ggt tcc acc tgc cac acc gtg tgc   2115
   Arg Cys Gly Val Cys His Gly Asn Gly Ser Thr Cys His Thr Val Ser
                   690                 695                 700

   ggg acc ttc gag gag gcc gag ggt ctg ggg tat gtg gat gtg ggg ctg   2163
55 Gly Thr Phe Glu Glu Ala Glu Gly Leu Gly Tyr Val Asp Val Gly Leu
               705                 710                 715

   atc tca gcg ggc gca cgc gag atc cgc atc caa gag ctt gcc gag gct   2211
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   acc acc ttc aca tac gca cgc agg ggc aac tgg gag aac ctc acg tcc   2355
 5 Thr Thr Phe Thr Tyr Ala Arg Arg Gly Asn Trp Glu Asn Leu Thr Ser
                 770                 775                 780

   ccg ggt ccc acc aag gag cct gtc tgg atc cag gtg cct gcc tcc cgt   2403
   Pro Gly Pro Thr Lys Glu Pro Val Trp Ile Gln Val Pro Ala Ser Arg
10             785                 790                 795

   ggc cca ggc ggg ggg agc aga ggc gga gtc ccc agg ccc agc acc ctc   2451
   Gly Pro Gly Gly Gly Ser Arg Gly Gly Val Pro Arg Pro Ser Thr Leu
         800                 805                 810
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   cat ggc agg tct cgt cct gga gga gtg agc cct ggt tca gtc aca gag   2499
   His Gly Arg Ser Arg Pro Gly Gly Val Ser Pro Gly Ser Val Thr Glu
       815                 820                 825

20 cct ggc tct gag cca ggc cct cct gct gcg gcc tct acc tca gtt tcc   2547
   Pro Gly Ser Glu Pro Gly Pro Pro Ala Ala Ala Ser Thr Ser Val Ser
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   cca tct tta aaa tgg ccc aat ctt gta gct gca gtt cac aga ggt ggc   2595
25 Pro Ser Leu Lys Trp Pro Asn Leu Val Ala Ala Val His Arg Gly Gly
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   tgg ggt caa gct cct tta gga ctg ggt gga tgg aga aga cac ttt gtg   2643
   Trp Gly Gln Ala Pro Leu Gly Leu Gly Gly Trp Arg Arg His Leu Val
30             865                 870                 875

   ctc atg ggc ccc cgc ctg ccc acc cag ctg ctg ttc cag gag agc aac   2691
   Leu Met Gly Pro Arg Leu Pro Thr Gln Leu Leu Phe Gln Glu Ser Asn
         880                 885                 890
35
   cct ggg gtg cac tac gag tac acc atc cac agg gag gca ggt ggc cac   2739
   Pro Gly Val His Tyr Glu Tyr Thr Ile His Arg Glu Ala Gly Gly His
       895                 900                 905

40 gac gag gtc ccg ccg ccc gtg ttc tcc tgg cat tat ggg ccc tgg acc   2787
   Asp Glu Val Pro Pro Pro Val Phe Ser Trp His Tyr Gly Pro Trp Thr
   910                 915                 920                 925

   aag tgc aca gtc acc tgc ggc aga ggt gag aag tgg ggc agg tac agc   2835
45 Lys Cys Thr Val Thr Cys Gly Arg Gly Glu Lys Trp Gly Arg Tyr Ser
                 930                 935                 940

   ccc acc tgc agg ggc tta gtg tct gga cag gga cac tgg ctt cag ctc   2883
   Pro Thr Cys Arg Gly Leu Val Ser Gly Gln Gly His Trp Leu Gln Leu
50             945                 950                 955

   cca gct cac tgc tgg gcc acc acg ggt ttg gaa gtt tgt ttc tct gag   2931
   Pro Ala His Cys Trp Ala Thr Thr Gly Leu Glu Val Cys Phe Ser Glu
         960                 965                 970
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   cct cag ttc tcc atc tgt gag atg agg cta gtg att gcc ctg tgt gcc   2979
   Pro Gln Phe Ser Ile Cys Glu Met Arg Leu Val Ile Ala Leu Cys Pro
       975                 980                 985

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   gatcagtcca gtggcctttc acacactgga ctttttccc taaattattt atcaaatgta ttta
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ctagt 3218

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Leu	Leu	Leu	Leu	Leu	Leu	Cys	Ala	Leu	Ala	Pro	Gly	Ala	Pro	Gly	Pro	Ala
			20					25					30		
Pro	Gly	Arg	Ala	Thr	Glu	Gly	Arg	Ala	Ala	Leu	Asp	Ile	Val	His	Pro
	35					40					45				
Val	Arg	Val	Asp	Ala	Gly	Gly	Ser	Phe	Leu	Ser	Tyr	Glu	Leu	Trp	Pro
50					55					60					
Arg	Ala	Leu	Arg	Lys	Arg	Asp	Val	Ser	Val	Arg	Arg	Asp	Ala	Pro	Ala
65				70					75					80	
Phe	Tyr	Glu	Leu	Gln	Tyr	Arg	Gly	Arg	Glu	Leu	Arg	Phe	Asn	Leu	Thr
			85					90					95		
Ala	Asn	Gln	His	Leu	Leu	Ala	Pro	Gly	Phe	Val	Ser	Glu	Thr	Arg	Arg
		100					105					110			
Arg	Gly	Gly	Leu	Gly	Arg	Ala	His	Ile	Arg	Ala	His	Thr	Pro	Ala	Cys
	115					120					125				
His	Leu	Leu	Gly	Glu	Val	Gln	Asp	Pro	Glu	Leu	Glu	Gly	Gly	Leu	Ala
130					135					140					
Ala	Ile	Ser	Ala	Cys	Asp	Gly	Leu	Lys	Gly	Val	Phe	Gln	Leu	Ser	Asn
145				150					155					160	
Glu	Asp	Tyr	Phe	Ile	Glu	Pro	Leu	Asp	Ser	Ala	Pro	Ala	Arg	Pro	Gly
			165					170					175		
His	Ala	Gln	Pro	His	Val	Val	Tyr	Lys	Arg	Gln	Ala	Pro	Glu	Arg	Leu
		180					185					190			
Ala	Gln	Arg	Gly	Asp	Ser	Ser	Ala	Pro	Ser	Thr	Cys	Gly	Val	Gln	Val
	195					200					205				
Tyr	Pro	Glu	Leu	Glu	Ser	Arg	Arg	Glu	Arg	Trp	Glu	Gln	Arg	Gln	Gln
210					215					220					
Trp	Arg	Arg	Pro	Arg	Leu	Arg	Arg	Leu	His	Gln	Arg	Ser	Val	Ser	Lys
225				230					235					240	
Glu	Leu	Tyr	Tyr	Glu	Gln	Leu	Val	Val	Ala	Asp	Ala	Leu	Met	Val	Glu
Met							Arg								
						200					205				

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Ile Thr Ile Val Arg Leu Val Leu Leu Glu Asp Glu Glu Glu Asp Leu
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Lys Ile Thr His His Ala Asp Asn Thr Leu Lys Ser Phe Cys Lys Trp
305                 310                 315                 320

Gln Lys Ser Ile Asn Met Lys Gly Asp Ala His Pro Leu His His Asp
                325                 330                 335

Thr Ala Ile Leu Leu Thr Arg Lys Asp Leu Cys Ala Ala Met Asn Arg
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Pro Cys Glu Thr Leu Gly Leu Ser His Val Ala Gly Met Cys Gln Pro
        355                 360                 365

His Arg Ser Cys Ser Ile Asn Glu Asp Thr Gly Leu Pro Leu Ala Phe
    370                 375                 380

Thr Val Ala His Glu Leu Gly His Ser Phe Gly Ile Gln His Asp Gly
385                 390                 395                 400

Ser Gly Asn Asp Cys Glu Pro Val Gly Lys Arg Pro Phe Ile Met Ser
                405                 410                 415

Pro Gln Leu Leu Tyr Asp Ala Ala Pro Leu Thr Trp Ser Arg Cys Ser
            420                 425                 430

Arg Gln Tyr Ile Thr Arg Phe Leu Asp Arg Gly Trp Gly Leu Cys Leu
        435                 440                 445

Asp Asp Pro Pro Ala Lys Asp Ile Ile Asp Phe Pro Ser Val Pro Pro
    450                 455                 460

Gly Val Leu Tyr Asp Val Ser His Gln Cys Arg Leu Gln Tyr Gly Ala
465                 470                 475                 480

Tyr Ser Ala Phe Cys Glu Asp Met Asp Asn Val Cys His Thr Leu Trp
                485                 490                 495

Cys Ser Val Gly Thr Thr Cys His Ser Lys Leu Asp Ala Ala Val Asp
            500                 505                 510

Gly Thr Arg Cys Gly Glu Asn Lys Trp Cys Leu Ser Gly Glu Cys Val
        515                 520                 525

Pro Val Gly Phe Arg Pro Glu Ala Val Asp Gly Gly Trp Ser Gly Trp
    530                 535                 540

Ser Ala Trp Ser Ile Cys Ser Arg Ser Cys Gly Met Gly Val Gln Ser
545                 550                 555                 560

Ala Glu Arg Gln Cys Thr Gln Pro Thr Pro Lys Tyr Lys Gly Arg Tyr
                565                 570                 575

Cys Val Gly Glu Arg Lys Arg Phe Arg Leu Cys Asn Leu Gln Ala Cys
            580                 585                 590

Pro Ala Gly Arg Pro Ser Phe Ala His Val Glu Cys Ser His Phe Asn
        595                 600                 605

Ala Ala Ala Ala Ala Ala Ala Ala Ala Ala Ala Ala Ala Ala Ala
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Ala Lys Lys Leu Arg Asp Ala Cys Val Asp Gly Thr Pro Cys Tyr Gln
645 650 655

Val Arg Ala Ser Arg Asp Leu Cys Ile Asn Gly Ile Cys Lys Asn Val
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Gly Cys Asp Phe Glu Ile Asp Ser Gly Ala Met Glu Asp Arg Cys Gly
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Val Cys His Gly Asn Gly Ser Thr Cys His Thr Val Ser Gly Thr Phe
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Trp Thr Ile Gln Trp Asn Gly Asp Tyr Gln Val Ala Gly Thr Thr Phe
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Thr Tyr Ala Arg Arg Gly Asn Trp Glu Asn Leu Thr Ser Pro Gly Pro
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Thr Lys Glu Pro Val Trp Ile Gln Val Pro Ala Ser Arg Gly Pro Gly
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Glu Pro Gly Pro Pro Ala Ala Ala Ser Thr Ser Val Ser Pro Ser Leu
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Pro Arg Leu Pro Thr Gln Leu Leu Phe Gln Glu Ser Asn Pro Gly Val
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His Tyr Gln Tyr Thr Ile His Arg Glu Ala Gly Gly His Asp Glu Val
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   Ser Glu Ala Arg Phe Val Glu Thr Leu Leu Val Ala Asp Ala Ser Met
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   gct gcc ttc tat ggg acc gac ctg cag aac cac atc ctc acg gtg atg  1063
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   tca atg gca gcc cga atc tac aag cac ccg agc atc agg aac tcc gtc  1111
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30 aac ctt gtg gtg gtg aaa gtg cta ata gtg gaa aaa gaa aga tgg ggc  1159
   Asn Leu Val Val Val Lys Val Leu Ile Val Glu Lys Glu Arg Trp Gly
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   cag tgt gac acc ctg ggg atg gca gac gtt ggc acc atc tgt gac ccc  1351
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   acc ctg gcc cat gag cta ggg cac gtt ctc agc atg ccc cat gat gat  1447
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25 Pro Gly His Leu Cys Leu Asp Gly Ser Cys Val Leu Lys Glu Asp Val
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   tca gga gtg tcc ccc cga gac cga tgc aag ctg ttt tgc aga gcc cgt   2215
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   ggg agg agt gag ttc aaa gtg ttt gaa gct aag gtg atc gat ggc act   2263
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   cgg ctg cag agc ttc cag gcc ctg cct gag cct ctt aca gta cag ctc   2688
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   Leu Thr Val Ser Gly Glu Val Phe Pro Pro Lys Val Arg Tyr Thr Phe
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   Ala Thr Thr Asn Ile Ile Gln Ser Leu Pro Ser Ala Glu Trp Val Leu
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Asn His Ile Leu Thr Leu Met Ser Val Ala Ala Arg Ile Tyr Lys His
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    65                  70                  75

tct gcc act gac ccc tgg cct gcc ttc gcc tcc tcc tct tcc tcc tct   287
Ser Ala Thr Asp Pro Trp Pro Ala Phe Ala Ser Ser Ser Ser Ser Ser
80                  85                  90                  95

acc tcc tcc cag gcg cat tac cgc ctc tct gcc ttc ggc cag cag ttt   335
Thr Ser Ser Gln Ala His Tyr Arg Leu Ser Ala Phe Gly Gln Gln Phe
                100                 105                 110

cta ttt aat ctc acc gcc aat gcc gga ttt atc gct cca ctg ttc act   383
Leu Phe Asn Leu Thr Ala Asn Ala Gly Phe Ile Ala Pro Leu Phe Thr
            115                 120                 125

gtc acc ctc ctt ggg acg ccc ggg gtg aat cag acc aag ttt tat tcc   431
Val Thr Leu Leu Gly Thr Pro Gly Val Asn Gln Thr Lys Phe Tyr Ser
        130                 135                 140

gaa gag gaa gcg gaa cta aag cac tgt ttc tac aaa agg cta tgt caa   479
Glu Glu Glu Ala Glu Leu Lys His Cys Phe Tyr Lys Arg Leu Cys Gln
    145                 150                 155

tac caa ctc cga gca cac ggc cgt cat cag cct ctg ctc agg aat gaa   527
Tyr Gln Leu Arg Ala His Gly Arg His Gln Pro Leu Leu Arg Asn Glu
160                 165                 170                 175

cac aaa aat agg cac agt aaa gac aag aag aaa acc aga gca aga aaa   575
His Lys Asn Arg His Ser Lys Asp Lys Lys Lys Thr Arg Ala Arg Lys

[illegible]                                                      71
Gly [illegible] Ala Leu Ala [illegible] Ala Tyr Gly Asn Lys Thr Asp Asn
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          210                 215                 220

   aca aga gaa aag agg acc cac aga agg aca aaa cgt ttt tta tcc tat   719
   Thr Arg Glu Lys Arg Thr His Arg Arg Thr Lys Arg Phe Leu Ser Tyr
 5     225                 230                 235

   cca cgg ttt gta gaa gtc ttg gtg gtg gca gac aac aga atg gtt tca   767
   Pro Arg Phe Val Glu Val Leu Val Val Ala Asp Asn Arg Met Val Ser
   240                 245                 250                 255
10
   tac cat gga gaa aac ctt caa cac tat att tta act tta atg tca att   815
   Tyr His Gly Glu Asn Leu Gln His Tyr Ile Leu Thr Leu Met Ser Ile
                   260                 265                 270

15 gta gcc tct atc tat aaa gac cca agt att gga aat tta att aat att   863
   Val Ala Ser Ile Tyr Lys Asp Pro Ser Ile Gly Asn Leu Ile Asn Ile
               275                 280                 285

   gtt att gtg aac tta att gtg att cat aat gaa cag gat ggg cct tcc   911
20 Val Ile Val Asn Leu Ile Val Ile His Asn Glu Gln Asp Gly Pro Ser
           290                 295                 300

   ata tct ttt aat gct cag aca aca tta aaa aac ttt tgt cag tgg cag   959
   Ile Ser Phe Asn Ala Gln Thr Thr Leu Lys Asn Phe Cys Gln Trp Gln
25     305                 310                 315

   cat tcg aac agt cca ggt gga atc cat cat gat act gct gtt ctc tta  1007
   His Ser Asn Ser Pro Gly Gly Ile His His Asp Thr Ala Val Leu Leu
   320                 325                 330                 335
30
   aca aga cag gat atc tgc aga gct cac gac aaa tgt gat acc tta ggc  1055
   Thr Arg Gln Asp Ile Cys Arg Ala His Asp Lys Cys Asp Thr Leu Gly
                   340                 345                 350

35 ctg gct gaa ctg gga acc att tgt gat ccc tat aga agt tgt tct att  1103
   Leu Ala Glu Leu Gly Thr Ile Cys Asp Pro Tyr Arg Ser Cys Ser Ile
               355                 360                 365

   agt gaa gat agt gga ttg agt aca gct ttt acg atc gct cat gag ctg  1151
40 Ser Glu Asp Ser Gly Leu Ser Thr Ala Phe Thr Ile Ala His Glu Leu
           370                 375                 380

   ggc cat gtg ttt aac atg cct cat gat gac aac aac aaa tgt aaa gaa  1199
   Gly His Val Phe Asn Met Pro His Asp Asp Asn Asn Lys Cys Lys Glu
45     385                 390                 395

   gaa gga gtt aag agt ccc cag cat gtc atg gct cca aca ctg aac ttc  1247
   Glu Gly Val Lys Ser Pro Gln His Val Met Ala Pro Thr Leu Asn Phe
   400                 405                 410                 415
50
   tac acc aac ccc tgg atg tgg tca aag tgt agt cga aaa tat atc act  1295
   Tyr Thr Asn Pro Trp Met Trp Ser Lys Cys Ser Arg Lys Tyr Ile Thr
                   420                 425                 430

55 gag ttt tta gac act ggt tat ggc gag tgt ttg ctt aac gaa cct gaa  1343
   Glu Phe Leu Asp Thr Gly Tyr Gly Glu Cys Leu Leu Asn Glu Pro Glu
               435                 440                 445

                                                                  1391
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    cca tat atg atg cag tgc aga cgg ctc tgg tgc aat aac gtc aat gga   1487
    Pro Tyr Met Met Gln Cys Arg Arg Leu Trp Cys Asn Asn Val Asn Gly
    480                 485                 490                 495

 5  gta cac aaa ggc tgc cgg act cag cac aca ccc tgg gcc gat ggg acg   1535
    Val His Lys Gly Cys Arg Thr Gln His Thr Pro Trp Ala Asp Gly Thr
                    500                 505                 510

    gag tgc gag cct gga aag cac tgc aag nat gga ttt tgt gtt ccc aaa   1583
10  Glu Cys Glu Pro Gly Lys His Cys Lys Xaa Gly Phe Cys Val Pro Lys
                515                 520                 525

    gaa atg gat gtc ccc gtg aca gat gga tcc tgg gga agt tgg agt ccc   1631
    Glu Met Asp Val Pro Val Thr Asp Gly Ser Trp Gly Ser Trp Ser Pro
15          530                 535                 540

    ttt gga acc tgc tcc aga aca tgt gga ggg ggc atc aaa aca gcc att   1679
    Phe Gly Thr Cys Ser Arg Thr Cys Gly Gly Gly Ile Lys Thr Ala Ile
        545                 550                 555
20
    cga gag tgc aac aga cca gaa cca aaa aat ggt gga aaa tac tgt gta   1727
    Arg Glu Cys Asn Arg Pro Glu Pro Lys Asn Gly Gly Lys Tyr Cys Val
    560                 565                 570                 575

25  gga cgt aga atg aaa ttt aag tcc tgc aac acg gag cca tgt ctc aag   1775
    Gly Arg Arg Met Lys Phe Lys Ser Cys Asn Thr Glu Pro Cys Leu Lys
                    580                 585                 590

    cag aag cga gac ttc cga gat gaa cag tgt gct cac ttt gac ggg aag   1823
30  Gln Lys Arg Asp Phe Arg Asp Glu Gln Cys Ala His Phe Asp Gly Lys
                595                 600                 605

    cat ttt aac atc aac ggt ctg ctt ccc aat gtg cgc tgg gtc cct aaa   1871
    His Phe Asn Ile Asn Gly Leu Leu Pro Asn Val Arg Trp Val Pro Lys
35          610                 615                 620

    tac agt gga att ctg atg aag gac cgg tgc aag ttg ttc tgc aga gtg   1919
    Tyr Ser Gly Ile Leu Met Lys Asp Arg Cys Lys Leu Phe Cys Arg Val
        625                 630                 635
40
    gca ggg aac aca gcc tac tat cag ctt cga gac aga gtg ata gat gga   1967
    Ala Gly Asn Thr Ala Tyr Tyr Gln Leu Arg Asp Arg Val Ile Asp Gly
    640                 645                 650                 655

45  act cct tgt ggc cag gac aca aat gat atc tgt gtc cag ggc ctt tgt   2015
    Thr Pro Cys Gly Gln Asp Thr Asn Asp Ile Cys Val Gln Gly Leu Cys
                    660                 665                 670

    cgg caa gct gga tgc gat cat gtt tta aac tca aaa gcc cgg aga gat   2063
50  Arg Gln Ala Gly Cys Asp His Val Leu Asn Ser Lys Ala Arg Arg Asp
                675                 680                 685

    aaa tgc ggg gtt tgt ggt ggc gat aat tct tca tgc aaa aca gtg gca   2111
    Lys Cys Gly Val Cys Gly Gly Asp Asn Ser Ser Cys Lys Thr Val Ala
55          690                 695                 700

    gga aca ttt aat aca gta cat tat ggt tac aat act gtg gtc cga att   2159
    Gly Thr Phe Asn Thr Val His Tyr Gly Tyr Asn Thr Val Val Arg Ile

    gaa aca gat gat gag aac tat tta gca tta tca gtt ttt gga tta aag   2207
    Glu Thr Asp Asp Glu Asn Tyr Leu Ala Leu Ser Val Phe Gly Leu Lys
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                    740                 745                 750

   ttc ttg cta aat gga aac ttt gtt gtc aca atg gcc aaa agg gaa att   2303
   Phe Leu Leu Asn Gly Asn Phe Val Val Thr Met Ala Lys Arg Glu Ile
 5             755                 760                 765

   cgc att ggg aat gct gtg gta gag tac agt ggg tcc gag act gcc gta   2351
   Arg Ile Gly Asn Ala Val Val Glu Tyr Ser Gly Ser Glu Thr Ala Val
            770                 775                 780
10
   gaa aga att aac tca aca gat cgc att gag caa gaa ctt ttg ctt cag   2399
   Glu Arg Ile Asn Ser Thr Asp Arg Ile Glu Gln Glu Leu Leu Leu Gln
        785                 790                 795

15 gtt ttg tcg gtg gga aag ttg tac aac ccc gat gta cgc tat tct ttc   2447
   Val Leu Ser Val Gly Lys Leu Tyr Asn Pro Asp Val Arg Tyr Ser Phe
   800                 805                 810                 815

   aat att cca att gaa gat aaa cct cag cag ttt tac tgg aac agt cat   2495
20 Asn Ile Pro Ile Glu Asp Lys Pro Gln Gln Phe Tyr Trp Asn Ser His
                   820                 825                 830

   ggg cca tgg caa gca tgc agt aaa ccc tgc caa ggg gaa cgg aaa cga   2543
   Gly Pro Trp Gln Ala Cys Ser Lys Pro Cys Gln Gly Glu Arg Lys Arg
25             835                 840                 845

   aaa ctt gtt tgc acc agg gaa tct gat cag ctt act gtt tct gat caa   2591
   Lys Leu Val Cys Thr Arg Glu Ser Asp Gln Leu Thr Val Ser Asp Gln
            850                 855                 860
30
   aga tgc gat cgg ctg ccc cag cct gga cac att act gaa ccc tgt ggt   2639
   Arg Cys Asp Arg Leu Pro Gln Pro Gly His Ile Thr Glu Pro Cys Gly
        865                 870                 875

35 aca ggc tgt gac ctg agg tgg cat gtt gcc agc agg agt gaa tgt agt   2687
   Thr Gly Cys Asp Leu Arg Trp His Val Ala Ser Arg Ser Glu Cys Ser
   880                 885                 890                 895

   gcc cag tgt ggc ttg ggt tac cgc aca ttg gac atc tac tgt gcc aaa   2735
40 Ala Gln Cys Gly Leu Gly Tyr Arg Thr Leu Asp Ile Tyr Cys Ala Lys
                   900                 905                 910

   tat agc agg ctg gat ggg aag act gag aag gtt gat gat ggt ttt tgc   2783
   Tyr Ser Arg Leu Asp Gly Lys Thr Glu Lys Val Asp Asp Gly Phe Cys
45             915                 920                 925

   agc agt cat ccc aaa cca agc aac cgt gaa aaa tgt tca ggg gaa tgt   2831
   Ser Ser His Pro Lys Pro Ser Asn Arg Glu Lys Cys Ser Gly Glu Cys
            930                 935                 940
50
   aac acg ggt ggc tgg cgc tat tct gcc tgg act gaa tgt tca aaa agc   2879
   Asn Thr Gly Gly Trp Arg Tyr Ser Ala Trp Thr Glu Cys Ser Lys Ser
        945                 950                 955

55 tgt gac ggt ggg acc cag agg aga agg gct att tgt gtc aat acc cga   2927
   Cys Asp Gly Gly Thr Gln Arg Arg Arg Ala Ile Cys Val Asn Thr Arg
   960                 965                 970                 975
```

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   tgg tca gag tgc ttg gtc act tgt gga aaa ggg cat aag cac agc cag   3071
   Trp Ser Glu Cys Leu Val Thr Cys Gly Lys Gly His Lys His Ser Gln
          1010                1015                1020

 5 gtc tgg tgt cag ttt ggt gaa gat cga tta aat gat aga atg tgt gac   3119
   Val Trp Cys Gln Phe Gly Glu Asp Arg Leu Asn Asp Arg Met Cys Asp
      1025                1030                1035

   cct gag acc aag cca aca tct atg cag act tgt cag cag ccg gaa tgt   3167
10 Pro Glu Thr Lys Pro Thr Ser Met Gln Thr Cys Gln Gln Pro Glu Cys
   1040                1045                1050                1055

   gca tcc tgg cag gcg ggt ccc tgg gta cag tgc agt gtc act tgt gga   3215
   Ala Ser Trp Gln Ala Gly Pro Trp Val Gln Cys Ser Val Thr Cys Gly
15                1060                1065                1070

   cag gga tac cag cta aga gca gtg aaa tgc atc att ggg act tat atg   3263
   Gln Gly Tyr Gln Leu Arg Ala Val Lys Cys Ile Ile Gly Thr Tyr Met
             1075                1080                1085
20
   tca gtg gta gat gac aat gac tgt aat gca gca act aga cca act gat   3311
   Ser Val Val Asp Asp Asn Asp Cys Asn Ala Ala Thr Arg Pro Thr Asp
          1090                1095                1100

25 acc cag gac tgt gaa tta cca tca tgt cat cct ccc cca gct gcc ccg   3359
   Thr Gln Asp Cys Glu Leu Pro Ser Cys His Pro Pro Pro Ala Ala Pro
      1105                1110                1115

   gaa acg agg aga agc aca tac agt gca cca aga acc cag tgg cga ttt   3407
30 Glu Thr Arg Arg Ser Thr Tyr Ser Ala Pro Arg Thr Gln Trp Arg Phe
   1120                1125                1130                1135

   ggg tct tgg acc cca tgc tca gcc act tgt ggg aaa ggt acc cgg atg   3455
   Gly Ser Trp Thr Pro Cys Ser Ala Thr Cys Gly Lys Gly Thr Arg Met
35                1140                1145                1150

   aga tac gtc agc tgc cga gat gag aat ggc tct gtg gct gac gag agt   3503
   Arg Tyr Val Ser Cys Arg Asp Glu Asn Gly Ser Val Ala Asp Glu Ser
             1155                1160                1165
40
   gcc tgt gct acc ctg cct aga cca gtg gca aag gaa gaa tgt tct gtg   3551
   Ala Cys Ala Thr Leu Pro Arg Pro Val Ala Lys Glu Glu Cys Ser Val
          1170                1175                1180

45 aca ccc tgt ggg caa tgg aag gcc ttg gac tgg agc tct tgc tct gtg   3599
   Thr Pro Cys Gly Gln Trp Lys Ala Leu Asp Trp Ser Ser Cys Ser Val
      1185                1190                1195

   acc tgt ggg caa ggt agg gca atc cgg caa gtg atg tgt gtc aac tac   3647
50 Thr Cys Gly Gln Gly Arg Ala Thr Arg Gln Val Met Cys Val Asn Tyr
   1200                1205                1210                1215

   agt gac cac gtg atc gat cgg agt gag tgt gac cag gat tat atc cca   3695
   Ser Asp His Val Ile Asp Arg Ser Glu Cys Asp Gln Asp Tyr Ile Pro
55                1220                1225                1230

   gaa act gac cag gac tgt tcc atg tca cca tgc cct caa agg acc cca   3743
   Glu Thr Asp Gln Asp Cys Ser Met Ser Pro Cys Pro Gln Arg Thr Pro

   [illegible]
   Arg Ser Ala Ser Pr  Ser Arg Thr His Val Leu Gly Gly Asn Gln Trp
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      1265                1270                1275

   aga act ggc ccc tgg gga gca tgt tcc agt acc tgt gct ggc gga tcc   3887
   Arg Thr Gly Pro Trp Gly Ala Cys Ser Ser Thr Cys Ala Gly Gly Ser
 5 1280                1285                1290                1295

   cag cgg cgt gtt gtt gta tgt cag gat gaa aat gga tac acc gca aac   3935
   Gln Arg Arg Val Val Val Cys Gln Asp Glu Asn Gly Tyr Thr Ala Asn
                  1300                1305                1310
10
   gac tgt gtg gag aga ata aaa cct gat gag caa aga gcc tgt gaa tcc   3983
   Asp Cys Val Glu Arg Ile Lys Pro Asp Glu Gln Arg Ala Cys Glu Ser
              1315                1320                1325

15 ggc cct tgt cct cag tgg gct tat ggc aac tgg gga gag tgc act aag   4031
   Gly Pro Cys Pro Gln Trp Ala Tyr Gly Asn Trp Gly Glu Cys Thr Lys
         1330                1335                1340

   ctg tgt ggt gga ggc ata aga aca aga ctg gtg gtc tct cag cgg tcc   4079
20 Leu Cys Gly Gly Gly Ile Arg Thr Arg Leu Val Val Ser Gln Arg Ser
      1345                1350                1355

   aac ggt gaa cgg ttt cca gat ttg agc tgt gaa att ctt gat aaa cct   4127
   Asn Gly Glu Arg Phe Pro Asp Leu Ser Cys Glu Ile Leu Asp Lys Pro
25 1360                1365                1370                1375

   ccc gat cgt gag cag tgt aac aca cat gct tgt cca cac gac gct gca   4175
   Pro Asp Arg Glu Gln Cys Asn Thr His Ala Cys Pro His Asp Ala Ala
                  1380                1385                1390
30
   tgg agt act ggc cct tgg agc tcg tgt tct gtc tct tgt ggt cga ggg   4223
   Trp Ser Thr Gly Pro Trp Ser Ser Cys Ser Val Ser Cys Gly Arg Gly
              1395                1400                1405

35 cat aaa caa cga aat gtt tac tgc atg gca aaa gat gga agc cat tta   4271
   His Lys Gln Arg Asn Val Tyr Cys Met Ala Lys Asp Gly Ser His Leu
         1410                1415                1420

   gaa agt gat tac tgt aag cac ctg gct aag cca cat ggg cac aga aag   4319
40 Glu Ser Asp Tyr Cys Lys His Leu Ala Lys Pro His Gly His Arg Lys
      1425                1430                1435

   tgc cga gga gga aga tgc ccc aaa tgg aaa gct ggc gct tgg agt cag   4367
   Cys Arg Gly Gly Arg Cys Pro Lys Trp Lys Ala Gly Ala Trp Ser Gln
45 1440                1445                1450                1455

   tgc tct gtg tcc atg ggc cga ggc gta cag cag agg cat gtg ggc tgt   4415
   Cys Ser Val Ser Met Gly Arg Gly Val Gln Gln Arg His Val Gly Cys
                  1460                1465                1470
50
   cag atc gga aca cac aaa ata gcc aga gag acc gag tgc aac cca tac   4463
   Gln Ile Gly Thr His Lys Ile Ala Arg Glu Thr Glu Cys Asn Pro Tyr
              1475                1480                1485

55 acc aga ccg gag tcg gaa tgc gaa tgc caa ggc cca cgg tgt ccc ctt   4511
   Thr Arg Pro Glu Ser Glu Cys Glu Cys Gln Gly Pro Arg Cys Pro Leu
         1490                1495                1500

   ... ... ... ... ... ... ... ... ... ... ... ... ... ... ... ...
   ... 1505                1510                1515                1520
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   gag gtg cat ggg gca cgc tgt gac gtg agc aag cgg ccg gtg gac cgt   4655
   Glu Val His Gly Ala Arg Cys Asp Val Ser Lys Arg Pro Val Asp Arg
                  1540                1545                1550

 5 gaa agc tgt agt ttg caa ccc tgc gag tat gtc tgg act aca gga gaa   4703
   Glu Ser Cys Ser Leu Gln Pro Cys Glu Tyr Val Trp Thr Thr Gly Glu
              1555                1560                1565

   tgg tca gag tgc tca gtg acc tgt gga aaa ggc tac aaa caa agg ctt   4751
10 Trp Ser Glu Cys Ser Val Thr Cys Gly Lys Gly Tyr Lys Gln Arg Leu
          1570                1575                1580

   gtc tcg tgc agc gag att tac acc ggg aaa gag aat tat gaa tac agc   4799
   Val Ser Cys Ser Glu Ile Tyr Thr Gly Lys Glu Asn Tyr Glu Tyr Ser
15    1585                1590                1595

   tac caa acc acc atc aac tgc cca ggc acg cag ccc ccc agt gtt cac   4847
   Tyr Gln Thr Thr Ile Asn Cys Pro Gly Thr Gln Pro Pro Ser Val His
   1600                1605                1610                1615
20
   ccc tgt tac ctg agg gag tgc cct gtc tcg gcc acc tgg aga gtt ggc   4895
   Pro Cys Tyr Leu Arg Glu Cys Pro Val Ser Ala Thr Trp Arg Val Gly
                  1620                1625                1630

25 aac tgg ggg agc tgc tca gtg tct tgt ggt gtt gga gtg atg cag aga   4943
   Asn Trp Gly Ser Cys Ser Val Ser Cys Gly Val Gly Val Met Gln Arg
              1635                1640                1645

   tct gtg caa tgt tta acc aat gag gac caa ccc agc cac tta tgc cac   4991
30 Ser Val Gln Cys Leu Thr Asn Glu Asp Gln Pro Ser His Leu Cys His
          1650                1655                1660

   act gat ctg aag cca gaa gaa cga aaa acc tgc cgt aat gtc tat aac   5039
   Thr Asp Leu Lys Pro Glu Glu Arg Lys Thr Cys Arg Asn Val Tyr Asn
35    1665                1670                1675

   tgt gag tta ccc cag aat tgc aag gag gta aaa aga ctt aaa ggt gcc   5087
   Cys Glu Leu Pro Gln Asn Cys Lys Glu Val Lys Arg Leu Lys Gly Ala
   1680                1685                1690                1695
40
   agt gaa gat ggt gaa tat ttc ctg atg att aga gga aag ctt ctg aag   5135
   Ser Glu Asp Gly Glu Tyr Phe Leu Met Ile Arg Gly Lys Leu Leu Lys
                  1700                1705                1710

45 ata ttc tgt gcg ggg atg cac tct gac cac ccc aaa gag tac gtg aca   5183
   Ile Phe Cys Ala Gly Met His Ser Asp His Pro Lys Glu Tyr Val Thr
              1715                1720                1725

   ctg gtg cat gga gac tct gag aat ttc tcc gag gtt tat ggg gac agg   5231
50 Leu Val His Gly Asp Ser Glu Asn Phe Ser Glu Val Tyr Gly His Arg
          1730                1735                1740

   tta cac aac cca aca gaa tgt ccc tat aac ggg agc cgg cgc gat gac   5279
   Leu His Asn Pro Thr Glu Cys Pro Tyr Asn Gly Ser Arg Arg Asp Asp
55    1745                1750                1755

   tgc caa tgt ggg aag gat tac agg gtc gct ggg ttt tcc agt ttt cag   5327
   Cys Gln Cys Asn Lys Asp Tyr Thr Ala Ala Gly Phe Ser Ser Phe Gln

   cag ttt gtc agg gca tgt ggg gga tat ... gtg ttt ctt gc ... ...   5...
   Gln Phe Ala Arg Thr Ser Glu Gly His Glu Val Pro Phe Ala Trp Ala
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             1795                1800                1805

   ggg gat tgc tac agc gct gcc aag tgc cca cag ggt cgt ttt agc atc   5471
   Gly Asp Cys Tyr Ser Ala Ala Lys Cys Pro Gln Gly Arg Phe Ser Ile
 5       1810                1815                1820

   aac ctt tat gga acc ggc ttg tct tta act gaa tct gcc aga tgg ata   5519
   Asn Leu Tyr Gly Thr Gly Leu Ser Leu Thr Glu Ser Ala Arg Trp Ile
      1825                1830                1835
10
   tca caa ggg aat tat gct gtc tct gac atc aag aag tcg ccg gat ggt   5567
   Ser Gln Gly Asn Tyr Ala Val Ser Asp Ile Lys Lys Ser Pro Asp Gly
   1840                1845                1850                1855

15 acc cga gtc gta ggg aaa tgc ggt ggt tac tgt gga aaa tgc act cca   5615
   Thr Arg Val Val Gly Lys Cys Gly Gly Tyr Cys Gly Lys Cys Thr Pro
                  1860                1865                1870

   tcc tct ggt act ggc ctg gag gtg cga gtt tta tagcttaggt gctttgaaga 5668
20 Ser Ser Gly Thr Gly Leu Glu Val Arg Val Leu
              1875                1880

   ggaagccatt atggatggat gaaggatagt aatgcaatac ctccacctta atttgggtgc 5728

25 atgtgtatgt gtgtgtgtgt ttgtgtgtga cttgtatgct tgtgtgtgta aatgtgtgta 5788

   catatacata tataca                                                 5804


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     1               5                  10                  15

   Arg Asp Leu Ala Glu Met Gly Ser Pro Asp Ala Ala Ala Ala Val Arg
40              20                  25                  30

   Lys Asp Arg Leu His Pro Arg Gln Val Lys Leu Leu Glu Thr Leu Ser
            35                  40                  45

45 Glu Tyr Glu Ile Val Ser Pro Ile Arg Val Asn Ala Leu Gly Glu Pro
        50                  55                  60

   Phe Pro Thr Asn Val His Phe Lys Arg Thr Arg Arg Ser Ile Asn Ser
    65                  70                  75                  80
90
   Ala Thr Asp Pro Trp Pro Ala Phe Ala Ser Ser Ser Ser Ser Ser Thr
                85                  90                  95

   Ser Ser Gln Ala His Tyr Arg Leu Ser Ala Phe Gly Gln Gln Phe Leu
55              100                 105                 110

   Phe Asn Leu Thr Ala Asn Ala Gly Phe Ile Ala Pro Leu Phe Thr Val
            115                 120                 125
```

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                                                    .
                                                    .
   Gln Leu Arg Ala His Gly Arg His Gln Pro Leu Leu Arg Asn Glu His
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                    165                 170                 175

   Lys Asn Arg His Ser Lys Asp Lys Lys Lys Thr Arg Ala Arg Lys Trp
               180                 185                 190

 5
   Gly Glu Arg Ile Asn Leu Ala Gly Asp Val Ala Ala Leu Asn Ser Gly
           195                 200                 205

   Leu Ala Thr Glu Ala Phe Ser Ala Tyr Gly Asn Lys Thr Asp Asn Thr
10     210                 215                 220

   Arg Glu Lys Arg Thr His Arg Arg Thr Lys Arg Phe Leu Ser Tyr Pro
   225                 230                 235                 240

15 Arg Phe Val Glu Val Leu Val Val Ala Asp Asn Arg Met Val Ser Tyr
                   245                 250                 255

   His Gly Glu Asn Leu Gln His Tyr Ile Leu Thr Leu Met Ser Ile Val
               260                 265                 270

20
   Ala Ser Ile Tyr Lys Asp Pro Ser Ile Gly Asn Leu Ile Asn Ile Val
           275                 280                 285

   Ile Val Asn Leu Ile Val Ile His Asn Glu Gln Asp Gly Pro Ser Ile
25     290                 295                 300

   Ser Phe Asn Ala Gln Thr Thr Leu Lys Asn Phe Cys Gln Trp Gln His
   305                 310                 315                 320

30 Ser Asn Ser Pro Gly Gly Ile His His Asp Thr Ala Val Leu Leu Thr
                   325                 330                 335

   Arg Gln Asp Ile Cys Arg Ala His Asp Lys Cys Asp Thr Leu Gly Leu
               340                 345                 350

35
   Ala Glu Leu Gly Thr Ile Cys Asp Pro Tyr Arg Ser Cys Ser Ile Ser
           355                 360                 365

   Glu Asp Ser Gly Leu Ser Thr Ala Phe Thr Ile Ala His Glu Leu Gly
40     370                 375                 380

   His Val Phe Asn Met Pro His Asp Asp Asn Asn Lys Cys Lys Glu Glu
   385                 390                 395                 400

45 Gly Val Lys Ser Pro Gln His Val Met Ala Pro Thr Leu Asn Phe Tyr
                   405                 410                 415

   Thr Asn Pro Trp Met Trp Ser Lys Cys Ser Arg Lys Tyr Ile Thr Glu
               420                 425                 430

50
   Phe Leu Asp Thr Gly Tyr Gly Glu Cys Leu Leu Asn Glu Pro Glu Ser
           435                 440                 445

   Arg Pro Tyr Pro Leu Pro Val Gln Leu Pro Gly Ile Leu Tyr Asn Val
55     450                 455                 460

   Asn Lys Gln Xaa Glu Leu Ile Phe Gly Pro Gly Ser Gln Val Cys Pro
   465                 470                 475                 480
```

```
   Lys Asn Arg Gly Lys His Cys Lys Asp Gly Asn Lys Val Pro Cys Ala
```

```
          515                 520                 525

   Met Asp Val Pro Val Thr Asp Gly Ser Trp Gly Ser Trp Ser Pro Phe
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   Gly Thr Cys Ser Arg Thr Cys Gly Gly Gly Ile Lys Thr Ala Ile Arg
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   Glu Cys Asn Arg Pro Glu Pro Lys Asn Gly Gly Lys Tyr Cys Val Gly
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   Arg Arg Met Lys Phe Lys Ser Cys Asn Thr Glu Pro Cys Leu Lys Gln
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15 Lys Arg Asp Phe Arg Asp Glu Gln Cys Ala His Phe Asp Gly Lys His
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   Phe Asn Ile Asn Gly Leu Leu Pro Asn Val Arg Trp Val Pro Lys Tyr
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   Gly Asn Thr Ala Tyr Tyr Gln Leu Arg Asp Arg Val Ile Asp Gly Thr
25                 645                 650                 655

   Pro Cys Gly Gln Asp Thr Asn Asp Ile Cys Val Gln Gly Leu Cys Arg
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30 Gln Ala Gly Cys Asp His Val Leu Asn Ser Lys Ala Arg Arg Asp Lys
          675                 680                 685

   Cys Gly Val Cys Gly Gly Asp Asn Ser Ser Cys Lys Thr Val Ala Gly
       690                 695                 700
35
   Thr Phe Asn Thr Val His Tyr Gly Tyr Asn Thr Val Val Arg Ile Pro
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   Ala Gly Ala Thr Asn Ile Asp Val Arg Gln His Ser Phe Ser Gly Glu
40                 725                 730                 735

   Thr Asp Asp Asp Asn Tyr Leu Ala Leu Ser Ser Ser Lys Gly Glu Phe
              740                 745                 750

45 Leu Leu Asn Gly Asn Phe Val Val Thr Met Ala Lys Arg Glu Ile Arg
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   Ile Gly Asn Ala Val Val Glu Tyr Ser Gly Ser Glu Thr Ala Val Glu
       770                 775                 780
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   Arg Ile Asn Ser Thr Asp Arg Ile Glu Gln Glu Leu Leu Leu Gln Val
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   Leu Ser Val Gly Lys Leu Tyr Asn Pro Asp Val Arg Tyr Ser Phe Asn
55                 805                 810                 815

   Ile Pro Ile Glu Asp Lys Pro Gln Gln Phe Tyr Trp Asn Ser His Leu
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   Leu Asp Arg Leu Ile Glu Asn Asp His Ile Thr Val Pro Tyr Asp Ile
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   Gly Cys Asp Leu Arg Trp His Val Ala Ser Arg Ser Glu Cys Ser Ala
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   Gln Cys Gly Leu Gly Tyr Arg Thr Leu Asp Ile Tyr Cys Ala Lys Tyr
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   Ser Arg Leu Asp Gly Lys Thr Glu Lys Val Asp Asp Gly Phe Cys Ser
10         915                 920                 925

   Ser His Pro Lys Pro Ser Asn Arg Glu Lys Cys Ser Gly Glu Cys Asn
       930                 935                 940

15 Thr Gly Gly Trp Arg Tyr Ser Ala Trp Thr Glu Cys Ser Lys Ser Cys
   945                 950                 955                 960

   Asp Gly Gly Thr Gln Arg Arg Arg Ala Ile Cys Val Asn Thr Arg Asn
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20
   Asp Val Leu Asp Asp Ser Lys Cys Thr His Gln Glu Lys Val Thr Ile
               980                 985                 990

   Gln Arg Cys Ser Glu Phe Pro Cys Pro Gln Trp Lys Ser Gly Asp Trp
25         995                1000                1005

   Ser Glu Cys Leu Val Thr Cys Gly Lys Gly His Lys His Ser Gln Val
      1010                1015                1020

30 Trp Cys Gln Phe Gly Glu Asp Arg Leu Asn Asp Arg Met Cys Asp Pro
   1025                1030                1035                1040

   Glu Thr Lys Pro Thr Ser Met Gln Thr Cys Gln Gln Pro Glu Cys Ala
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35
   Ser Trp Gln Ala Gly Pro Trp Val Gln Cys Ser Val Thr Cys Gly Gln
              1060                1065                1070

   Gly Tyr Gln Leu Arg Ala Val Lys Cys Ile Ile Gly Thr Tyr Met Ser
40        1075                1080                1085

   Val Val Asp Asp Asn Asp Cys Asn Ala Ala Thr Arg Pro Thr Asp Thr
      1090                1095                1100

45 Gln Asp Cys Glu Leu Pro Ser Cys His Pro Pro Pro Ala Ala Pro Glu
   1105                1110                1115                1120

   Thr Arg Arg Ser Thr Tyr Ser Ala Pro Arg Thr Gln Trp Arg Pro Gly
                  1125                1130                1135

50
   Ser Trp Thr Pro Cys Ser Ala Thr Cys Gly Lys Gly Thr Arg Met Arg
              1140                1145                1150

   Tyr Val Ser Cys Arg Asp Glu Asn Gly Ser Val Ala Asp Glu Ser Ala
55        1155                1160                1165

   Cys Ala Thr Leu Pro Arg Pro Val Ala Lys Glu Glu Cys Ser Val Thr
      1170                1175                1180

                  ..                  ...                 ...

   Asp His Val Ile Arg Arg Ser Glu Cys Asp Gln Asp Tyr Lys Pro Gly
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   Thr Asp Gln Asp Cys Ser Met Ser Pro Cys Pro Gln Arg Thr Pro Asp
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 5
   Ser Gly Leu Ala Gln His Pro Phe Gln Asn Glu Asp Tyr Arg Pro Arg
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   Ser Ala Ser Pro Ser Arg Thr His Val Leu Gly Gly Asn Gln Trp Arg
10 1265                1270                1275                1280

   Thr Gly Pro Trp Gly Ala Cys Ser Ser Thr Cys Ala Gly Gly Ser Gln
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15 Arg Arg Val Val Val Cys Gln Asp Glu Asn Gly Tyr Thr Ala Asn Asp
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   Cys Val Glu Arg Ile Lys Pro Asp Glu Gln Arg Ala Cys Glu Ser Gly
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20
   Pro Cys Pro Gln Trp Ala Tyr Gly Asn Trp Gly Glu Cys Thr Lys Leu
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   Cys Gly Gly Gly Ile Arg Thr Arg Leu Val Val Ser Gln Arg Ser Asn
25 1345                1350                1355                1360

   Gly Glu Arg Phe Pro Asp Leu Ser Cys Glu Ile Leu Asp Lys Pro Pro
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30 Asp Arg Glu Gln Cys Asn Thr His Ala Cys Pro His Asp Ala Ala Trp
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   Ser Thr Gly Pro Trp Ser Ser Cys Ser Val Ser Cys Gly Arg Gly His
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35
   Lys Gln Arg Asn Val Tyr Cys Met Ala Lys Asp Gly Ser His Leu Glu
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   Ser Asp Tyr Cys Lys His Leu Ala Lys Pro His Gly His Arg Lys Cys
40 1425                1430                1435                1440

   Arg Gly Gly Arg Cys Pro Lys Trp Lys Ala Gly Ala Trp Ser Gln Cys
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45 Ser Val Ser Met Gly Arg Gly Val Gln Gln Arg His Val Gly Cys Gln
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   Ile Gly Thr His Lys Ile Ala Arg Glu Thr Glu Cys Asn Pro Tyr Thr
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50
   Arg Pro Glu Ser Glu Cys Glu Cys Gln Gly Pro Arg Cys Pro Leu Tyr
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   Thr Trp Arg Ala Glu Glu Trp Gln Glu Cys Thr Lys Thr Cys Gly Glu
55 1505                1510                1515                1520

   Gly Ser Arg Tyr Arg Lys Val Val Cys Val Asp Asp Asn Lys Asn Glu
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   Ala Ala Lys Ser Val Thr Lys Arg Lys Pro Tyr Lys Gln Arg Leu Val
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      1570                1575                1580

   Ser Cys Ser Glu Ile Tyr Thr Gly Lys Glu Asn Tyr Glu Tyr Ser Tyr
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 5
   Gln Thr Thr Ile Asn Cys Pro Gly Thr Gln Pro Pro Ser Val His Pro
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   Cys Tyr Leu Arg Glu Cys Pro Val Ser Ala Thr Trp Arg Val Gly Asn
10            1620                1625                1630

   Trp Gly Ser Cys Ser Val Ser Cys Gly Val Gly Val Met Gln Arg Ser
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15 Val Gln Cys Leu Thr Asn Glu Asp Gln Pro Ser His Leu Cys His Thr
      1650                1655                1660

   Asp Leu Lys Pro Glu Glu Arg Lys Thr Cys Arg Asn Val Tyr Asn Cys
   1665                1670                1675                1680
20
   Glu Leu Pro Gln Asn Cys Lys Glu Val Lys Arg Leu Lys Gly Ala Ser
                   1685                1690                1695

   Glu Asp Gly Glu Tyr Phe Leu Met Ile Arg Gly Lys Leu Leu Lys Ile
25            1700                1705                1710

   Phe Cys Ala Gly Met His Ser Asp His Pro Lys Glu Tyr Val Thr Leu
          1715                1720                1725

30 Val His Gly Asp Ser Glu Asn Phe Ser Glu Val Tyr Gly His Arg Leu
      1730                1735                1740

   His Asn Pro Thr Glu Cys Pro Tyr Asn Gly Ser Arg Arg Asp Asp Cys
   1745                1750                1755                1760
35
   Gln Cys Arg Lys Asp Tyr Thr Ala Ala Gly Phe Ser Ser Phe Gln Lys
                   1765                1770                1775

   Ile Arg Ile Asp Leu Thr Ser Met Gln Ile Ile Thr Thr Asp Leu Gln
40            1780                1785                1790

   Phe Ala Arg Thr Ser Glu Gly His Pro Val Pro Phe Ala Thr Ala Gly
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45 Asp Cys Tyr Ser Ala Ala Lys Cys Pro Gln Gly Arg Phe Ser Ile Asn
      1810                1815                1820

   Leu Tyr Gly Thr Gly Leu Ser Leu Thr Glu Ser Ala Arg Trp Ile Ser
   1825                1830                1835                1840
50
   Gln Gly Asn Tyr Ala Val Ser Asp Ile Lys Lys Ser Pro Asp Gly Thr
                   1845                1850                1855

   Arg Val Val Gly Lys Cys Gly Gly Tyr Cys Gly Lys Cys Thr Pro Ser
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tct cac gat gga gat tat ttc att gaa cca ctg cag tct gtg gat gag 97
Ser His Asp Gly Asp Tyr Phe Ile Glu Pro Leu Gln Ser Val Asp Glu
20 25 30

caa gag gat gaa gag gaa caa aac aaa ccc cac att att tat agg cac 145
Gln Glu Asp Glu Glu Glu Gln Asn Lys Pro His Ile Ile Tyr Arg His
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agc acc cct cag agg gaa ccc tcc aca gga aag cat gcc tgt gcc acc 193
Ser Thr Pro Gln Arg Glu Pro Ser Thr Gly Lys His Ala Cys Ala Thr
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tca gaa ctc aaa aat agt cac agt aaa gac aag cgg aaa atc aga atg 241
Ser Glu Leu Lys Asn Ser His Ser Lys Asp Lys Arg Lys Ile Arg Met
65 70 75 80

cga aaa cgg aga aag agg aat agc ctg gct gac gac gtg gca ctg cta 289
Arg Lys Arg Arg Lys Arg Asn Ser Leu Ala Asp Asp Val Ala Leu Leu
85 90 95

aag agc ggt ttg gca aca aag gtg ctc tct ggc tat agc aac cag aca 337
Lys Ser Gly Leu Ala Thr Lys Val Leu Ser Gly Tyr Ser Asn Gln Thr
100 105 110

aac aac aca agg gac aga tgg aac cac aaa aga acc aaa cgc ttt ctg 385
Asn Asn Thr Arg Asp Arg Trp Asn His Lys Arg Thr Lys Arg Phe Leu
115 120 125

tcc tac cca cgg ttt gta gag gtg atg gtg gtg gct gac cac agg atg 433
Ser Tyr Pro Arg Phe Val Glu Val Met Val Val Ala Asp His Arg Met
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gtt tta tac cac gga gca aac ctt caa cat tat atc tta acc tta atg 481
Val Leu Tyr His Gly Ala Asn Leu Gln His Tyr Ile Leu Thr Leu Met
145 150 155 160

tcc att gta gct tct atc tat aaa gac tca agt att gga aat tta att 529
Ser Ile Val Ala Ser Ile Tyr Lys Asp Ser Ser Ile Gly Asn Leu Ile
165 170 175

aat att gtt att gtg aac tta gtt gtg att cat aat gaa cag gag gga 577
Asn Ile Val Ile Val Asn Leu Val Val Ile His Asn Glu Gln Glu Gly
180 185 190

cct tac ata aat ttc aat gcc cag aca aca tta aag aac ttt tgc cag 625
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   tgt tcc att agt gaa gac agt ggg ctg agc aca gct ttc aca ata gct   817
 5 Cys Ser Ile Ser Glu Asp Ser Gly Leu Ser Thr Ala Phe Thr Ile Ala
               260                 265                 270

   cac gag ctg ggc cat gtg ttt aat atg cct cac gat gac agc aat aaa   865
   His Glu Leu Gly His Val Phe Asn Met Pro His Asp Asp Ser Asn Lys
10         275                 280                 285

   tgc aaa gaa gaa gga gtt aag agt ccc cag cat gtc atg gca cca aca   913
   Cys Lys Glu Glu Gly Val Lys Ser Pro Gln His Val Met Ala Pro Thr
       290                 295                 300
15
   ctg aac ttc tac acc aac ccc tgg atg tgg tca aag tgc agt cgg aaa   961
   Leu Asn Phe Tyr Thr Asn Pro Trp Met Trp Ser Lys Cys Ser Arg Lys
   305                 310                 315                 320

20 tac atc act gag ttc cta gac act ggg tac gga gag tgc ttg ctg aat  1009
   Tyr Ile Thr Glu Phe Leu Asp Thr Gly Tyr Gly Glu Cys Leu Leu Asn
                   325                 330                 335

   gaa cct gca tcc agg acc tat cct ttg cct tcc caa ctg ccc ggc ctt  1057
25 Glu Pro Ala Ser Arg Thr Tyr Pro Leu Pro Ser Gln Leu Pro Gly Leu
               340                 345                 350

   ctc tac aac gtg aat aaa caa tgt gaa ctg att ttt ggg cca ggc tct  1105
   Leu Tyr Asn Val Asn Lys Gln Cys Glu Leu Ile Phe Gly Pro Gly Ser
30         355                 360                 365

   caa gtg tgc ccc tat atg atg cag tgc aga cgg ctc tgg tgc aat aat  1153
   Gln Val Cys Pro Tyr Met Met Gln Cys Arg Arg Leu Trp Cys Asn Asn
       370                 375                 380
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   gtg gat gga gca cac aaa ggc tgc aag act cag cac acg ccc tgg gca  1201
   Val Asp Gly Ala His Lys Gly Cys Lys Thr Gln His Thr Pro Trp Ala
   385                 390                 395                 400

40 gat gga acc gag tgt gag cct gga aag cac tgc aag ttt gga ttt tgt  1249
   Asp Gly Thr Glu Cys Glu Pro Gly Lys His Cys Lys Phe Gly Phe Cys
                   405                 410                 415

   gtt ccc aaa gaa atg gag gcc cct gca att gat gga tcc tgg gga ggt  1297
45 Val Pro Lys Glu Met Glu Gly Pro Ala Ile Asp Gly Ser Trp Gly Gly
               420                 425                 430

   tgg agc cac ttt ggg acc tgc tca aga acg tgt gga gga ggc att aaa  1345
   Trp Ser His Phe Gly Thr Cys Ser Arg Thr Cys Gly Gly Gly Ile Lys
50         435                 440                 445

   aca gcc atc aga gag tgc aac aga cca gag cca aaa aat ggt ggg aag  1393
   Thr Ala Ile Arg Glu Cys Asn Arg Pro Glu Pro Lys Asn Gly Gly Lys
       450                 455                 460
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   tac tgt gta gga agg aga atg aag ttc aaa tcc tgc aac acg gag ccc  1441
   Tyr Cys Val Gly Arg Arg Met Lys Phe Lys Ser Cys Asn Thr Glu Pro
   465                 470                 475                 480

   Asp Lys Cys His Pro Asn Ile Asn Gly Val Lys Arg Trp
                   485                 490
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   ttt cct aag tac agc gga att ttg atg aag gac cgg tgc aag ttg ttc   1585
   Phe Pro Lys Tyr Ser Gly Ile Leu Met Lys Asp Arg Cys Lys Leu Phe
           515                 520                 525
5
   tgc aga gtg gca gga aac aca gcc tac tac cag ctc cga gac aga gtg   1633
   Cys Arg Val Ala Gly Asn Thr Ala Tyr Tyr Gln Leu Arg Asp Arg Val
       530                 535                 540

10 att gac gga acc cct tgt ggc cag gac aca aat gac atc tgt gtc caa   1681
   Ile Asp Gly Thr Pro Cys Gly Gln Asp Thr Asn Asp Ile Cys Val Gln
   545                 550                 555                 560

   ggc ctt tgc cgg caa gct gga tgt gat cat att tta aac tca aag gtc   1729
15 Gly Leu Cys Arg Gln Ala Gly Cys Asp His Ile Leu Asn Ser Lys Val
                   565                 570                 575

   cgg aaa gat aaa tgt ggg att tgt ggt gga gat aat tct tca tgc aaa   1777
   Arg Lys Asp Lys Cys Gly Ile Cys Gly Gly Asp Asn Ser Ser Cys Lys
20             580                 585                 590

   aca gtg gca gga aca ttt aac act gtc cat tat ggt tac aat act gtt   1825
   Thr Val Ala Gly Thr Phe Asn Thr Val His Tyr Gly Tyr Asn Thr Val
           595                 600                 605
25
   gtc cga att ccg gct ggt gct acc agc att gac gtg cgt cag cac agc   1873
   Val Arg Ile Pro Ala Gly Ala Thr Ser Ile Asp Val Arg Gln His Ser
       610                 615                 620

30 ttc tca ggg aag tct gag gat gac aac tac cta gct tta tca aac agt   1921
   Phe Ser Gly Lys Ser Glu Asp Asp Asn Tyr Leu Ala Leu Ser Asn Ser
   625                 630                 635                 640

   aaa ggt gaa ttc ctg cta aat gga gac ttt gtt gtc tcc atg tcc aaa   1969
35 Lys Gly Glu Phe Leu Leu Asn Gly Asp Phe Val Val Ser Met Ser Lys
                   645                 650                 655

   agg gag gtc cgc gtg ggg agc gcc gtc att gag tac agc gga tcg gac   2017
   Arg Glu Val Arg Val Gly Ser Ala Val Ile Glu Tyr Ser Gly Ser Asp
40             660                 665                 670

   aat gtg tgt gaa aga ctg aac tgt acg gac cgt atc gag gaa gaa ctt   2065
   Asn Val Cys Glu Arg Leu Asn Cys Thr Asp Arg Ile Glu Glu Glu Leu
           675                 680                 685
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   ctc ctt cag gtg ttg tcc gtg gga aag ctg tat aac cca gat gtg cgg   2113
   Leu Leu Gln Val Leu Ser Val Gly Lys Leu Tyr Asn Pro Asp Val Arg
       690                 695                 700

50 tac tca ttc aac att ccc att gag gac aaa cct cag cag ttt tac tgg   2161
   Tyr Ser Phe Asn Ile Pro Ile Glu Asp Lys Pro Gln Gln Phe Tyr Trp
   705                 710                 715                 720

   aac agt cac ggg ccg tgg caa gca tgc agc aag ccc tgc caa ggg gag   2209
55 Asn Ser His Gly Pro Trp Gln Ala Cys Ser Lys Pro Cys Gln Gly Glu
                   725                 730                 735

   ggc agg cga aaa ctt gtt tgc acc aag gat gaa tct gat ggc ttg agt   2257
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   ggg tgt agt gaa gaa gtt gaa ttg agg tgg tat gtt gat gga aag agg   2305
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   Ala Cys Gly Thr Asp Cys Asp Leu Arg Trp His Val Ala Ser Lys Ser
       770                 775                 780

   gaa tgc agt gcc cag tgt ggt ttg ggc tac cgt act tta gac atc cac   2401
 5 Glu Cys Ser Ala Gln Cys Gly Leu Gly Tyr Arg Thr Leu Asp Ile His
   785                 790                 795                 800

   tgt gcc aaa tac agc agg atg gac ggg aag acg gag aag gtg gat gac   2449
   Cys Ala Lys Tyr Ser Arg Met Asp Gly Lys Thr Glu Lys Val Asp Asp
10                 805                 810                 815

   agt ttc tgt agc agt caa ccc aga ccg agt aac cag gag aaa tgc tca   2497
   Ser Phe Cys Ser Ser Gln Pro Arg Pro Ser Asn Gln Glu Lys Cys Ser
              820                 825                 830
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   gga gag tgc agc aca ggt gga tgg cgc tat tca gcc tgg acc gaa tgt   2545
   Gly Glu Cys Ser Thr Gly Gly Trp Arg Tyr Ser Ala Trp Thr Glu Cys
          835                 840                 845

20 tct aga agc tgt gat ggt ggt acc cac aga aga aga gca att tgt gtc   2593
   Ser Arg Ser Cys Asp Gly Gly Thr His Arg Arg Arg Ala Ile Cys Val
       850                 855                 860

   aac acc cgc aat gat gtc ctg gat gac agc aa                       2625
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   Gln Glu Asp Glu Glu Glu Gln Asn Lys Pro His Ile Ile Tyr Arg His
          35                  40                  45

   Ser Thr Pro Gln Arg Glu Pro Ser Thr Gly Lys His Ala Cys Ala Thr
45      50                  55                  60

   Ser Glu Leu Lys Asn Ser His Ser Lys Asp Lys Arg Lys Ile Arg Met
    65                  70                  75                  80

50 Arg Lys Arg Arg Lys Arg Asn Ser Leu Ala Asp Asp Val Ala Leu Leu
                   85                  90                  95

   Lys Ser Gly Leu Ala Thr Lys Val Leu Ser Gly Tyr Ser Asn Gln Thr
              100                 105                 110
55
   Asn Asn Thr Arg Asp Arg Trp Asn His Lys Arg Thr Lys Arg Phe Leu
          115                 120                 125

   Thr Ile Val Ala Arg Tyr Lys Arg Thr Val Ile Arg Asn Leu Ile
                  135                 140                 145
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Asn Ile Val Ile Val Asn Leu Val Val Ile His Asn Glu Gln Glu Gly
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Pro Tyr Ile Asn Phe Asn Ala Gln Thr Thr Leu Lys Asn Phe Cys Gln
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Trp Gln His Ser Lys Asn Tyr Leu Gly Gly Ile Gln His Asp Thr Ala
    210                 215                 220

Val Leu Val Thr Arg Glu Asp Ile Cys Arg Ala Gln Asp Lys Cys Asp
225                 230                 235                 240

Thr Leu Gly Leu Ala Glu Leu Gly Thr Ile Cys Asp Pro Tyr Arg Ser
                245                 250                 255

Cys Ser Ile Ser Glu Asp Ser Gly Leu Ser Thr Ala Phe Thr Ile Ala
            260                 265                 270

His Glu Leu Gly His Val Phe Asn Met Pro His Asp Asp Ser Asn Lys
        275                 280                 285

Cys Lys Glu Glu Gly Val Lys Ser Pro Gln His Val Met Ala Pro Thr
    290                 295                 300

Leu Asn Phe Tyr Thr Asn Pro Trp Met Trp Ser Lys Cys Ser Arg Lys
305                 310                 315                 320

Tyr Ile Thr Glu Phe Leu Asp Thr Gly Tyr Gly Glu Cys Leu Leu Asn
                325                 330                 335

Glu Pro Ala Ser Arg Thr Tyr Pro Leu Pro Ser Gln Leu Pro Gly Leu
            340                 345                 350

Leu Tyr Asn Val Asn Lys Gln Cys Glu Leu Ile Phe Gly Pro Gly Ser
        355                 360                 365

Gln Val Cys Pro Tyr Met Met Gln Cys Arg Arg Leu Trp Cys Asn Asn
    370                 375                 380

Val Asp Gly Ala His Lys Gly Cys Lys Thr Gln His Thr Pro Trp Ala
385                 390                 395                 400

Asp Gly Thr Glu Cys Glu Pro Gly Lys His Cys Lys Phe Gly Phe Cys
                405                 410                 415

Val Pro Lys Glu Met Glu Gly Pro Ala Ile Asp Gly Ser Trp Gly Gly
            420                 425                 430

Trp Ser His Phe Gly Thr Cys Ser Arg Thr Cys Gly Gly Gly Ile Lys
        435                 440                 445

Thr Ala Ile Arg Glu Cys Asn Arg Pro Glu Pro Lys Asn Gly Gly Lys
    450                 455                 460

Tyr Cys Val Gly Arg Arg Met Lys Phe Lys Ser Cys Asn Thr Glu Pro
465                 470                 475                 480

Ile Pro Lys Tyr Met Gly Leu Leu Met Tyr Asp Arg Cys Tyr Leu Phe
                485                 490                 495
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Cys Arg Val Ala Gly Asn Thr Ala Tyr Tyr Gln Leu Arg Asp Arg Val
530 535 540

Ile Asp Gly Thr Pro Cys Gly Gln Asp Thr Asn Asp Ile Cys Val Gln
545 550 555 560

Gly Leu Cys Arg Gln Ala Gly Cys Asp His Ile Leu Asn Ser Lys Val
565 570 575

Arg Lys Asp Lys Cys Gly Ile Cys Gly Gly Asp Asn Ser Ser Cys Lys
580 585 590

Thr Val Ala Gly Thr Phe Asn Thr Val His Tyr Gly Tyr Asn Thr Val
595 600 605

Val Arg Ile Pro Ala Gly Ala Thr Ser Ile Asp Val Arg Gln His Ser
610 615 620

Phe Ser Gly Lys Ser Glu Asp Asp Asn Tyr Leu Ala Leu Ser Asn Ser
625 630 635 640

Lys Gly Glu Phe Leu Leu Asn Gly Asp Phe Val Val Ser Met Ser Lys
645 650 655

Arg Glu Val Arg Val Gly Ser Ala Val Ile Glu Tyr Ser Gly Ser Asp
660 665 670

Asn Val Cys Glu Arg Leu Asn Cys Thr Asp Arg Ile Glu Glu Glu Leu
675 680 685

Leu Leu Gln Val Leu Ser Val Gly Lys Leu Tyr Asn Pro Asp Val Arg
690 695 700

Tyr Ser Phe Asn Ile Pro Ile Glu Asp Lys Pro Gln Gln Phe Tyr Trp
705 710 715 720

Asn Ser His Gly Pro Trp Gln Ala Cys Ser Lys Pro Cys Gln Gly Glu
725 730 735

Arg Arg Pro Lys Leu Val Cys Thr Arg Glu Ser Asp Gln Leu Thr Val
740 745 750

Ser Asp Gln Arg Cys Asp Arg Leu Pro Gln Pro Gly Pro Val Thr Glu
755 760 765

Ala Cys Gly Thr Asp Cys Asp Leu Arg Trp His Val Ala Ser Lys Ser
770 775 780

Glu Cys Ser Ala Gln Cys Gly Leu Gly Tyr Arg Thr Leu Asp Ile His
785 790 795 800

Cys Ala Lys Tyr Ser Arg Met Asp Gly Lys Thr Glu Lys Val Asp Asp
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Asn Thr Ala Asn Asp Val Lys Asp Asp
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gct atg aga tcg cct tcc tca ccc gcg tgg acc aca acg ggg cac tgc   96
Ala Met Arg Ser Pro Ser Pro Pro Ala Trp Thr Thr Thr Gly His Cys
             20                  25                  30

tgg cct tct cgc cac ctc ctc ccc gga gca gcg ccg cgg cac ggg ggc  144
Trp Pro Ser Arg His Leu Leu Pro Gly Ala Ala Pro Arg His Gly Gly
         35                  40                  45

cac agc cga gtc ccg cct ctt cta caa agt ggc ctc gcc agc acc cac  192
His Ser Arg Val Pro Pro Leu Leu Gln Ser Gly Leu Ala Ser Thr His
     50                  55                  60

ttc ctg ctg aac ctg acc cgc agc tcc cgt cta ctg gca ggg cgc gtc  240
Phe Leu Leu Asn Leu Thr Arg Ser Ser Arg Leu Leu Ala Gly Arg Val
 65                  70                  75                  80

tcc gtg gag tac tgg aca cgg gag ggc ctg gcc tgg cag agg gcg gcc  288
Ser Val Glu Tyr Trp Thr Arg Glu Gly Leu Ala Trp Gln Arg Ala Ala
                 85                  90                  95

cgg ccc cac tgc ctc tac gct ggt cac ctg cag ggc cag gcc agc agc  336
Arg Pro His Cys Leu Tyr Ala Gly His Leu Gln Gly Gln Ala Ser Ser
            100                 105                 110

tcc cat gtg gcc atc agc acc tgt gga ggc ctg cac ggc ctg atc gtg  384
Ser His Val Ala Ile Ser Thr Cys Gly Gly Leu His Gly Leu Ile Val
        115                 120                 125

gca gac gag gaa gag tac ctg att gag ccc ctg cac ggt ggg ccc aag  432
Ala Asp Glu Glu Glu Tyr Leu Ile Glu Pro Leu His Gly Gly Pro Lys
    130                 135                 140

ggt tct cgg agt ccg gag gaa agt gga cca cat tgt gtg tac aag cgt  480
Gly Ser Arg Ser Pro Glu Glu Ser Gly Pro His Cys Val Tyr Lys Arg
145                 150                 155                 160

tcc tcc ctg cgt cac ccc cac ctg gac aca gcc tgt gga gtg aga gat  528
Ser Ser Leu Arg His Pro His Leu Asp Thr Ala Cys Gly Val Arg Asp
                165                 170                 175

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       210                 215                 220

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 5 Ala Asp Lys Met Met Val Ala Tyr His Gly Arg Arg Asp Val Glu Gln
   225                 230                 235                 240

   tat gtc ctg gcc atc atg aac att gtt gcc aaa ctt ttc cag gac tcg   768
   Tyr Val Leu Ala Ile Met Asn Ile Val Ala Lys Leu Phe Gln Asp Ser
10                 245                 250                 255

   agt ctg gga agc acc gtt aac atc ctc gta act cgc ctc atc ctg ctc   816
   Ser Leu Gly Ser Thr Val Asn Ile Leu Val Thr Arg Leu Ile Leu Leu
               260                 265                 270
15
   acg gag gac cag ccc act ctg gag atc acc cac cat gcc ggg aag tcc   864
   Thr Glu Asp Gln Pro Thr Leu Glu Ile Thr His His Ala Gly Lys Ser
           275                 280                 285

20 cta gac agc ttc tgt aag tgg cag aaa tcc atc gtg aac cac agc ggc   912
   Leu Asp Ser Phe Cys Lys Trp Gln Lys Ser Ile Val Asn His Ser Gly
       290                 295                 300

   cat ggc aat gcc att cca gag aac ggt gtg gct aac cat gac aca gca   960
25 His Gly Asn Ala Ile Pro Glu Asn Gly Val Ala Asn His Asp Thr Ala
   305                 310                 315                 320

   gtg ctc atc aca cgc tat gac atc tgc atc tac aag aac aaa ccc tgc  1008
   Val Leu Ile Thr Arg Tyr Asp Ile Cys Ile Tyr Lys Asn Lys Pro Cys
30                 325                 330                 335

   ggc aca cta ggc ctg gcc cgg tgg gcg gaa tgt gtg agc gcg aga gaa  1056
   Gly Thr Leu Gly Leu Ala Arg Trp Ala Glu Cys Val Ser Ala Arg Glu
               340                 345                 350
35
   gct gca gcg tca atg agg aca ttg gct gcc aca agc gtt cac cat tgc  1104
   Ala Ala Ala Ser Met Arg Thr Leu Ala Ala Thr Ser Val His His Cys
           355                 360                 365

40 cac gag atc ggg cac aca ttc ggc atg aac cat gac ggc gtg gga aac  1152
   His Glu Ile Gly His Thr Phe Gly Met Asn His Asp Gly Val Gly Asn
       370                 375                 380

   agc tgt ggg gcc cgt ggt cag gac cca gcc aag ctc atg gct gcc cac  1200
45 Ser Cys Gly Ala Arg Gly Gln Asp Pro Ala Lys Leu Met Ala Ala His
   385                 390                 395                 400

   att acc atg aag acc aac cca ttc gtg tgg tca tcc tgc aac cgt gac  1248
   Ile Thr Met Lys Thr Asn Pro Phe Val Trp Ser Ser Cys Asn Arg Asp
50                 405                 410                 415

   tac atc acc agc ttt cta gac tcg ggc ctg ggg ctc tgc ctg aac aac  1296
   Tyr Ile Thr Ser Phe Leu Asp Ser Gly Leu Gly Leu Cys Leu Asn Asn
               420                 425                 430
55
   ggg ccc ccg aga cag gac ttt gtg tac ccg acg gtg gca ccg ggc caa  1344
   Arg Pro Pro Arg Gln Asp Phe Val Tyr Pro Thr Val Ala Pro Gly Gln
           435                 440                 445

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   445                 450                 455
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   aag agc aac cgg tgc atc acc aac agc atc ccg gcc gcc gag ggc acg   1488
   Lys Ser Asn Arg Cys Ile Thr Asn Ser Ile Pro Ala Ala Glu Gly Thr
                   485                 490                 495
5
   ctg tgc cag acg cac acc atc gac aag ggg tgg tgc tac aaa cgg gtc   1536
   Leu Cys Gln Thr His Thr Ile Asp Lys Gly Trp Cys Tyr Lys Arg Val
               500                 505                 510

10 tgt gtc ccc ttt ggg tcg cgc cca gag ggt gtg gac gga gcc tgg ggg   1584
   Cys Val Pro Phe Gly Ser Arg Pro Glu Gly Val Asp Gly Ala Trp Gly
           515                 520                 525

   ccg tgg act cca tgg ggc gac tgc agc cgg acc tgt ggc ggc ggc gtg   1632
15 Pro Trp Thr Pro Trp Gly Asp Cys Ser Arg Thr Cys Gly Gly Gly Val
       530                 535                 540

   tcc tct tct agt cgt cac tgc gac agc ccc agg cca acc atc ggg ggc   1680
   Ser Ser Ser Ser Arg His Cys Asp Ser Pro Arg Pro Thr Ile Gly Gly
20 545                 550                 555                 560

   aag tac tgt ctg ggt gag aga agg cgg cac cgc tcc tgc aac acg gat   1728
   Lys Tyr Cys Leu Gly Glu Arg Arg Arg His Arg Ser Cys Asn Thr Asp
                   565                 570                 575
25
   gac tgt ccc cct ggc tcc cag gac ttc aga gaa gtg cag tgt gct gaa   1776
   Asp Cys Pro Pro Gly Ser Gln Asp Phe Arg Glu Val Gln Cys Ala Glu
               580                 585                 590

30 ttt gac agc atc cct ttc cgt ggg aaa ttc tac aag tgg aaa acg tac   1824
   Phe Asp Ser Ile Pro Phe Arg Gly Lys Phe Tyr Lys Trp Lys Thr Tyr
           595                 600                 605

   cgg gga ggg ggc gtg aag gcc tgc tcg ctc acg agc cta gtg gaa ggc   1872
35 Arg Gly Gly Gly Val Lys Ala Cys Ser Leu Thr Ser Leu Ala Glu Gly
       610                 615                 620

   ttc aac ttc tac acg gag agg gcg gca gcc gtg gtg gac ggg aca ccc   1920
   Phe Asn Phe Tyr Thr Glu Arg Ala Ala Ala Val Val Asp Gly Thr Pro
40 625                 630                 635                 640

   tgc cgt cca gac acg gtg gac att tgc gtc agt ggc gaa tgc aag cac   1968
   Cys Arg Pro Asp Thr Val Asp Ile Cys Val Ser Gly Glu Cys Lys His
                   645                 650                 655
45
   gtg ggc tgc gac cga gtc ctg ggc tcc gac ctg cgg gag gac aag tgc   2016
   Val Gly Cys Asp Arg Val Leu Gly Ser Asp Leu Arg Glu Asp Lys Cys
               660                 665                 670

   agg gtg tgt ggc ggt gac ggc agt gcc tgc gag acc atc gag ggc gtc   2064
   Arg Val Cys Gly Gly Asp Gly Ser Ala Cys Glu Thr Ile Glu Gly Val
           675                 680                 685

   ttc agc cca gcc tca cct ggg gcc ggg tac gag gat gtc gtc tgg att   2112
55 Phe Ser Pro Ala Ser Pro Gly Ala Gly Tyr Glu Asp Val Val Trp Ile
       690                 695                 700

   ccc aaa ggc tgc gtg gac atg ttc atc cag gat ctg aac gtc ttt gtc   2160

   gtg ctt ggg agc ccc aac ctg gag gtg ctc gta gat ggg ttt aac
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   Leu Pro Gly Thr Pro Gln Pro His Arg Leu Pro Leu Ala Gly Thr Thr
                740                 745                 750

   ttt caa ctg cga cag ggg cca gac cag gtc cag agc ctc gaa gcc ctg   2304
 5 Phe Gln Leu Arg Gln Gly Pro Asp Gln Val Gln Ser Leu Glu Ala Leu
          755                 760                 765

   gga ccg att aat gca tct ctc atc gtc atg gtg ctg gcc cgg acc gag   2352
   Gly Pro Ile Asn Ala Ser Leu Ile Val Met Val Leu Ala Arg Thr Glu
10     770                 775                 780

   ctg cct gcc ctc cgc tac cgc ttc aat gcc ccc atc gcc cgt gac tcg   2400
   Leu Pro Ala Leu Arg Tyr Arg Phe Asn Ala Pro Ile Ala Arg Asp Ser
   785                 790                 795                 800
15
   ctg ccc ccc tac tcc tgg cac tat gcg ccc tgg acc aag tgc tcg gcc   2448
   Leu Pro Pro Tyr Ser Trp His Tyr Ala Pro Trp Thr Lys Cys Ser Ala
                805                 810                 815

20 cag tgt gca ggc ggt agc cag gtg cag gcg gtg gag tgc cgc aac cag   2496
   Gln Cys Ala Gly Gly Ser Gln Val Gln Ala Val Glu Cys Arg Asn Gln
                820                 825                 830

   ctg gac agc tcc gcg gtc gcc ccc cac tac tgc agt gcc cac agc aag   2544
25 Leu Asp Ser Ser Ala Val Ala Pro His Tyr Cys Ser Ala His Ser Lys
          835                 840                 845

   ctg ccc aaa agg cag cgc gcc tgc aac acg gag cct tgc cct cca gac   2592
   Leu Pro Lys Arg Gln Arg Ala Cys Asn Thr Glu Pro Cys Pro Pro Asp
30     850                 855                 860

   tgg gtt gta ggg aac tgg tcg ctc tgc agc cgc agc tgc gat gca ggc   2640
   Trp Val Val Gly Asn Trp Ser Leu Cys Ser Arg Ser Cys Asp Ala Gly
   865                 870                 875                 880
35
   gtg cgc agt acg tcg gtc gtg tgc cag cgc cgc gtc tct gcc gcg gag   2688
   Val Arg Ser Thr Ser Val Val Cys Gln Arg Arg Val Ser Ala Ala Glu
                885                 890                 895

40 gag aag gcg ctg gac gac agc gca tgc ccg cag ccg cgc cca cct gta   2736
   Glu Lys Ala Leu Asp Asp Ser Ala Cys Pro Gln Pro Arg Pro Pro Val
                900                 905                 910

   ctg gag gcc tgc cac ggt ccc act tgc cct ccg gag tgg gca acc ctc   2784
45 Leu Glu Ala Cys His Gly Pro Thr Cys Pro Pro Glu Trp Ala Thr Leu
          915                 920                 925

   gac tgg tct gag tgt acc cca agc tgt ggg cct ggt ctc cgc cac cga   2832
   Asp Trp Ser Glu Cys Thr Pro Ser Cys Gly Pro Gly Leu Arg His Arg
50     930                 935                 940

   gtg gtc ctt tgt aag agt gca gat caa cga tct act ctg ccc cct ggg   2880
   Val Val Leu Cys Lys Ser Ala Asp Gln Arg Ser Thr Leu Pro Pro Gly
   945                 950                 955                 960
55
   cac tgc ctt cct gca gcc aag cca cca tct act atg cga tgt aac ttg   2928
   His Cys Leu Pro Ala Ala Lys Pro Pro Ser Thr Met Arg Cys Asn Leu
                965                 970                 975

   [illegible]                                                      [illegible]
01 Ser Thr Gln Lys Arg Leu Gly Ala Leu Asn Ser Val Arg Lys Thr
          980                 985                 990
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Ser His Thr Gly Gln Pro Ser Arg Glu Cys Thr Glu Ala Leu Arg Pro
1010 1015 1020

tcc acc atg cag cag tgt gag gcc aag tgt gac agt gtg gtg ccg cct 3120
Ser Thr Met Gln Gln Cys Glu Ala Lys Cys Asp Ser Val Val Pro Pro
1025 1030 1035 1040

gga gat ggc cca gaa gaa tgc aag gat gtg aac aag gtg gct tac tgc 3168
Gly Asp Gly Pro Glu Glu Cys Lys Asp Val Asn Lys Val Ala Tyr Cys
1045 1050 1055

ccc ctg gtg ctc aaa ttt cag ttc tgt agc cga gcc tac ttc cgc cag 3216
Pro Leu Val Leu Lys Phe Gln Phe Cys Ser Arg Ala Tyr Phe Arg Gln
1060 1065 1070

atg tgc tgc aaa acc tgc caa ggc cgc tag ggtacctgga accaacctgg 3266
Met Cys Cys Lys Thr Cys Gln Gly Arg
1075 1080

agtacaggct gaggcagggg acatcccact ggagagggca tgagggaaag gggggcttga 3326

attgaagggt gagatgcagt tgaaagttat ttattgggta accctacagg gctcctgact 3386

aaggggtgga gaagagctgg ctacccaggg accctctgct gtatcttgcc cagttgatag 3446

tgaagagaga ggactccttg ttgcacacat atttaagtcc ctagcacccc tcccaccctt 3506

tgatcggaat atgtactgtg aagagtgggg gtggggaggg gtgtgctggt gccctgcccc 3566

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aggcttgata ccacctccct gtagccctcc cccagactga cgaaggggaa gatccacccc 3686

aacctctgcc ctgcctgccc caggggggag ttcaacatcc aggccgttcc ccatcatggt 3746

gctacaagcc ctgccctggg gcccacacac tcctcaccaa gaagccttac attaaaaaag 3806

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Trp Pro Ser Arg His Leu Leu Pro Gly Ala Ala Pro Arg His Gly Gly
35 40 45
His Ser Arg Val Pro Pro Leu Leu Gln Ser Gly Leu Ala Ser Thr His
50 55 60

Val Ala Ile His Thr Gly Arg Trp Leu Leu Asp Leu Leu
115 120 125

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   Ala Asp Glu Glu Glu Tyr Leu Ile Glu Pro Leu His Gly Gly Pro Lys
       130                 135                 140
   Gly Ser Arg Ser Pro Glu Glu Ser Gly Pro His Cys Val Tyr Lys Arg
   145                 150                 155                 160
 5 Ser Ser Leu Arg His Pro His Leu Asp Thr Ala Cys Gly Val Arg Asp
                   165                 170                 175
   Glu Lys Pro Trp Lys Gly Arg Pro Trp Trp Leu Arg Thr Leu Lys Pro
               180                 185                 190
   Pro Pro Ala Arg Pro Leu Gly Asn Glu Thr Glu Arg Gly Gln Pro Gly
10         195                 200                 205
   Leu Lys Arg Ser Val Ser Arg Glu Arg Tyr Val Glu Thr Met Asp Val
       210                 215                 220
   Ala Asp Lys Met Met Val Ala Tyr His Gly Arg Arg Asp Val Glu Gln
   225                 230                 235                 240
15 Tyr Val Leu Ala Ile Met Asn Ile Val Ala Lys Leu Phe Gln Asp Ser
                   245                 250                 255
   Ser Leu Gly Ser Thr Val Asn Ile Leu Val Thr Arg Leu Ile Leu Leu
               260                 265                 270
   Thr Glu Asp Gln Pro Thr Leu Glu Ile Thr His His Ala Gly Lys Ser
20         275                 280                 285
   Leu Asp Ser Phe Cys Lys Trp Gln Lys Ser Ile Val Asn His Ser Gly
       290                 295                 300
   His Gly Asn Ala Ile Pro Glu Asn Gly Val Ala Asn His Asp Thr Ala
   305                 310                 315                 320
25 Val Leu Ile Thr Arg Tyr Asp Ile Cys Ile Tyr Lys Asn Lys Pro Cys
                   325                 330                 335
   Gly Thr Leu Gly Leu Ala Arg Trp Ala Glu Cys Val Ser Ala Arg Glu
               340                 345                 350
   Ala Ala Ala Ser Met Arg Thr Leu Ala Ala Thr Ser Val His His Cys
30         355                 360                 365
   His Glu Ile Gly His Thr Phe Gly Met Asn His Asp Gly Val Gly Asn
       370                 375                 380
   Ser Cys Gly Ala Arg Gly Gln Asp Pro Ala Lys Leu Met Ala Ala His
   385                 390                 395                 400
35 Ile Thr Met Lys Thr Asn Pro Phe Val Trp Ser Ser Cys Asn Arg Asp
                   405                 410                 415
   Tyr Ile Thr Ser Phe Leu Asp Ser Gly Leu Gly Leu Cys Leu Asn Asn
               420                 425                 430
   Arg Pro Pro Arg Gln Asp Phe Val Tyr Pro Thr Val Ala Pro Gly Gln
40         435                 440                 445
   Ala Tyr Asp Ala Asp Glu Gln Cys Arg Phe Gln His Gly Val Lys Ser
       450                 455                 460
   Arg Gln Cys Lys Tyr Gly Glu Val Cys Ser Glu Leu Trp Cys Leu Ser
   465                 470                 475                 480
45 Lys Ser Asn Arg Cys Ile Thr Asn Ser Ile Pro Ala Ala Glu Gly Thr
                   485                 490                 495
   Leu Cys Gln Thr His Thr Ile Asp Lys Gly Trp Cys Tyr Lys Arg Val
               500                 505                 510
   Cys Val Pro Phe Gly Ser Arg Pro Glu Gly Val Asp Gly Ala Trp Gly
50         515                 520                 525
   Pro Trp Thr Pro Trp Gly Asp Cys Ser Arg Thr Cys Gly Gly Gly Val
       530                 535                 540
   Ser Ser Ser Ser Arg His Cys Asp Ser Pro Arg Pro Thr Ile Gly Gly
   545                 550                 555                 560
55 Lys Tyr Cys Leu Gly Glu Arg Arg Arg His Arg Ser Cys Asn Thr Asp
                   565                 570                 575
   Asp Cys Pro Pro Gly Ser Gln Asp Phe Arg Glu Val Gln Cys Ala Glu
               580                 585                 590

Val Tyr Arg Asp Ile Val Asp Leu Tyr Val Met Gly Leu Cys Lys His
           595                 600                 605
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Val Gly Cys Asp Arg Val Leu Gly Ser Asp Leu Arg Glu Asp Lys Cys
            660                 665                 670
Arg Val Cys Gly Gly Asp Gly Ser Ala Cys Glu Thr Ile Glu Gly Val
        675                 680                 685
Phe Ser Pro Ala Ser Pro Gly Ala Gly Tyr Glu Asp Val Val Trp Ile
    690                 695                 700
Pro Lys Gly Ser Val His Ile Phe Ile Gln Asp Leu Asn Leu Ser Leu
705                 710                 715                 720
Ser His Leu Ala Leu Lys Gly Asp Gln Glu Ser Leu Leu Leu Glu Gly
                725                 730                 735
Leu Pro Gly Thr Pro Gln Pro His Arg Leu Pro Leu Ala Gly Thr Thr
            740                 745                 750
Phe Gln Leu Arg Gln Gly Pro Asp Gln Val Gln Ser Leu Glu Ala Leu
        755                 760                 765
Gly Pro Ile Asn Ala Ser Leu Ile Val Met Val Leu Ala Arg Thr Glu
    770                 775                 780
Leu Pro Ala Leu Arg Tyr Arg Phe Asn Ala Pro Ile Ala Arg Asp Ser
785                 790                 795                 800
Leu Pro Pro Tyr Ser Trp His Tyr Ala Pro Trp Thr Lys Cys Ser Ala
                805                 810                 815
Gln Cys Ala Gly Gly Ser Gln Val Gln Ala Val Glu Cys Arg Asn Gln
            820                 825                 830
Leu Asp Ser Ser Ala Val Ala Pro His Tyr Cys Ser Ala His Ser Lys
        835                 840                 845
Leu Pro Lys Arg Gln Arg Ala Cys Asn Thr Glu Pro Cys Pro Pro Asp
    850                 855                 860
Trp Val Val Gly Asn Trp Ser Leu Cys Ser Arg Ser Cys Asp Ala Gly
865                 870                 875                 880
Val Arg Ser Thr Ser Val Val Cys Gln Arg Arg Val Ser Ala Ala Glu
                885                 890                 895
Glu Lys Ala Leu Asp Asp Ser Ala Cys Pro Gln Pro Arg Pro Pro Val
            900                 905                 910
Leu Glu Ala Cys His Gly Pro Thr Cys Pro Pro Glu Trp Ala Thr Leu
        915                 920                 925
Asp Trp Ser Glu Cys Thr Pro Ser Cys Gly Pro Gly Leu Arg His Arg
    930                 935                 940
Val Val Leu Cys Lys Ser Ala Asp Gln Arg Ser Thr Leu Pro Pro Gly
945                 950                 955                 960
His Cys Leu Pro Ala Ala Lys Pro Pro Ser Thr Met Arg Cys Asn Leu
                965                 970                 975
Arg Arg Cys Pro Pro Ala Arg Trp Val Thr Ser Glu Trp Gly Glu Cys
            980                 985                 990
Ser Thr Gln Cys Gly Leu Gly Gln Gln Gln Arg Thr Val Arg Cys Thr
        995                1000                1005
Ser His Thr Gly Gln Pro Ser Arg Glu Cys Thr Glu Ala Leu Arg Pro
    1010                1015                1020
Ser Thr Met Gln Gln Cys Glu Ala Lys Cys Asp Ser Val Val Pro Pro
1025                1030                1035                1040
Gly Asp Gly Pro Glu Glu Cys Lys Asp Val Asn Lys Val Ala Tyr Cys
                1045                1050                1055
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Ile Cys Val Ser Gly Glu Cys Lys His Val Gly Cys Asp Arg Leu Leu
            20                  25                  30

ggt tct gat ctc cga gag gac aaa tgc cgt gtg tgt ggg ggt gat ggc   145
Gly Ser Asp Leu Arg Glu Asp Lys Cys Arg Val Cys Gly Gly Asp Gly
        35                  40                  45

agt gcc tgt gag acc att gaa ggt gtc ttt agc cca gct ttg cca gga   193
Ser Ala Cys Glu Thr Ile Glu Gly Val Phe Ser Pro Ala Leu Pro Gly
    50                  55                  60

act ggg tat gag gac gtc gtc tgg atc ccc aaa ggc tcg gtc cac att   241
Thr Gly Tyr Glu Asp Val Val Trp Ile Pro Lys Gly Ser Val His Ile
65                  70                  75                  80

ttc atc caa gat ctg aac ctg tcc ctg agt cac ctg gcc cta aag ggg   289
Phe Ile Gln Asp Leu Asn Leu Ser Leu Ser His Leu Ala Leu Lys Gly
                85                  90                  95

gac caa gag tct ctg cta ctg gag ggg cta cct ggg acc ccc caa cct   337
Asp Gln Glu Ser Leu Leu Leu Glu Gly Leu Pro Gly Thr Pro Gln Pro
            100                 105                 110

nac cgc ctt ccc ctg gnt ggg acc aca ttt cat cta cgg cag ggg ccg   385
Xaa Arg Leu Pro Leu Xaa Gly Thr Thr Phe His Leu Arg Gln Gly Pro
        115                 120                 125

gac cag gca cag agc ctg gaa gcc ctg gga ccc att aat gca tct ctc   433
Asp Gln Ala Gln Ser Leu Glu Ala Leu Gly Pro Ile Asn Ala Ser Leu
    130                 135                 140

ttc atc atg gtg ctg gcc cag gca gag ttg cct gct ctc cac tac cgc   481
Ile Ile Met Val Leu Ala Gln Ala Glu Leu Pro Ala Leu His Tyr Arg
145                 150                 155                 160

ttc aat gca ccc att gcc cgg gat gca ctg ccc ccc tac tcc tgg cac   529
Phe Asn Ala Pro Ile Ala Arg Asp Ala Leu Pro Pro Tyr Ser Trp His
                165                 170                 175

tat gcc ccc tgg acc aaa tgc tca gcc cag tgt gca ggc ggc agc cag   577
Tyr Ala Pro Trp Thr Lys Cys Ser Ala Gln Cys Ala Gly Gly Ser Gln
            180                 185                 190

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   225                 230                 235                 240

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 5 Arg Cys Ser Arg Ser Cys Asp Ala Gly Val Arg Ser Arg Ser Val Val
                   245                 250                 255

   tgc caa cgc cgg gtg tct gct gca gag gaa aaa gcc tta gac gac agt   817
   Cys Gln Arg Arg Val Ser Ala Ala Glu Glu Lys Ala Leu Asp Asp Ser
10             260                 265                 270

   gcc tgt cca cag cca cgc cca cct gtg ctg gag gcc tgc caa ggc cca   865
   Ala Cys Pro Gln Pro Arg Pro Pro Val Leu Glu Ala Cys Gln Gly Pro
         275                 280                 285
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   atg tgc cct cct gag tgg gca acc ctg gac tgg tct gag tgt acc cca   913
   Met Cys Pro Pro Glu Trp Ala Thr Leu Asp Trp Ser Glu Cys Thr Pro
     290                 295                 300

20 agc tgt ggg cct ggt ctc cgc cac cga gtg gtc ctt tgt aag agt gca   961
   Ser Cys Gly Pro Gly Leu Arg His Arg Val Val Leu Cys Lys Ser Ala
   305                 310                 315                 320

   gat caa cga tct act ctg ccc cct ggg cac tgc ctt cct gca gcc aag  1009
25 Asp Gln Arg Ser Thr Leu Pro Pro Gly His Cys Leu Pro Ala Ala Lys
                   325                 330                 335

   cca cca tct act atg cga tgt aac ttg cgc cgc tgc cct cct gcc cgc  1057
   Pro Pro Ser Thr Met Arg Cys Asn Leu Arg Arg Cys Pro Pro Ala Arg
30             340                 345                 350

   tgg gtg acc agt gag tgg ggt gag tgt tcc aca cag tgt ggc ctc ggc  1105
   Trp Val Thr Ser Glu Trp Gly Glu Cys Ser Thr Gln Cys Gly Leu Gly
         355                 360                 365
35
   cag cag cag cgc aca gtg cgc tgc acc agc cac acc ggc cag cca tct  1153
   Gln Gln Gln Arg Thr Val Arg Cys Thr Ser His Thr Gly Gln Pro Ser
     370                 375                 380

40 cga gag tgc act gaa gcc ttg cgg cca tcc acc atg cag cag tgt gag  1201
   Arg Glu Cys Thr Glu Ala Leu Arg Pro Ser Thr Met Gln Gln Cys Glu
   385                 390                 395                 400

   gcc aag tgt gac agt gtg gtg ccg cct gga gat ggc cca gaa gaa tgc  1249
45 Ala Lys Cys Asp Ser Val Val Pro Pro Gly Asp Gly Pro Glu Glu Cys
                   405                 410                 415

   aag gat gtg aac aag gtg gct tac tgc ccc ctg gtg ctc aaa ttt cag  1297
   Lys Asp Val Asn Lys Val Ala Tyr Cys Pro Leu Val Leu Lys Phe Gln
50             420                 425                 430

   ttc tgt agc cga gcc tac ttc cgc cag atg agc tgc aaa acc tgc caa  1345
   Phe Cys Ser Arg Ala Tyr Phe Arg Gln Met Ser Cys Lys Thr Cys Gln
         435                 440                 445
55
   ggc cgc tagggtacct ggaaatcaacc tggagcacag gttgaggcag gggacatccc  1401
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35 40 45

Ser Ala Cys Glu Thr Ile Glu Gly Val Phe Ser Pro Ala Leu Pro Gly
50 55 60

Thr Gly Tyr Glu Asp Val Val Trp Ile Pro Lys Gly Ser Val His Ile
65 70 75 80

Phe Ile Gln Asp Leu Asn Leu Ser Leu Ser His Leu Ala Leu Lys Gly
85 90 95

Asp Gln Glu Ser Leu Leu Leu Glu Gly Leu Pro Gly Thr Pro Gln Pro
100 105 110

Xaa Arg Leu Pro Leu Xaa Gly Thr Thr Phe His Leu Arg Gln Gly Pro
115 120 125

Asp Gln Ala Gln Ser Leu Glu Ala Leu Gly Pro Ile Asn Ala Ser Leu
130 135 140

Ile Ile Met Val Leu Ala Gln Ala Glu Leu Pro Ala Leu His Tyr Arg
145 150 155 160

Phe Asn Ala Pro Ile Ala Arg Asp Ala Leu Pro Pro Tyr Ser Trp His
165 170 175

Tyr Ala Pro Trp Thr Lys Cys Ser Ala Gln Cys Ala Gly Gly Ser Gln
180 185 190

Val Gln Val Val Glu Cys Arg Asn Gln Leu Asp Ser Ser Ala Val Ala
195 200 205

Pro His Tyr Lys Ser Gly His Ser Cys Leu Pro Lys Arg Gln Arg Ala
210 215 220

Cys Asn Thr Glu Pro Cys Pro Pro Asp Trp Val Val Gly Asn Trp Ser
225 230 235 240

Arg Cys Ser Arg Ser Cys Asp Ala Gly Val Arg Ser Arg Ser Val Val
245 250 255

Met Lys Ile Glu Met Ala Asp Gly Ala Leu Cys Asp Thr Leu Glu His
260 265 270

```
   Ser Cys Gly Pro Gly Leu Arg His Arg Val Val Leu Cys Lys Ser Ala
   305                 310                 315                 320

 5 Asp Gln Arg Ser Thr Leu Pro Pro Gly His Cys Leu Pro Ala Ala Lys
                   325                 330                 335

   Pro Pro Ser Thr Met Arg Cys Asn Leu Arg Arg Cys Pro Pro Ala Arg
               340                 345                 350
10
   Trp Val Thr Ser Glu Trp Gly Glu Cys Ser Thr Gln Cys Gly Leu Gly
           355                 360                 365

   Gln Gln Gln Arg Thr Val Arg Cys Thr Ser His Thr Gly Gln Pro Ser
15     370                 375                 380

   Arg Glu Cys Thr Glu Ala Leu Arg Pro Ser Thr Met Gln Gln Cys Glu
   385                 390                 395                 400

20 Ala Lys Cys Asp Ser Val Val Pro Pro Gly Asp Gly Pro Glu Glu Cys
                   405                 410                 415

   Lys Asp Val Asn Lys Val Ala Tyr Cys Pro Leu Val Leu Lys Phe Gln
               420                 425                 430
25
   Phe Cys Ser Arg Ala Tyr Phe Arg Gln Met Ser Cys Lys Thr Cys Gln
           435                 440                 445

   Gly Arg
30     450



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45                                                          Met Glu
                                                             1

   tgc tgc cgt cgg gca act cct ggg aca ctg ctc ctc ttt ctg gct ttc   104
   Cys Cys Arg Arg Ala Thr Pro Gly Thr Leu Leu Leu Phe Leu Ala Phe
50           5                  10                  15

   ctg ctc ctg agt tcc agg acc gca cgc tcc gag gag gac cgg gac ggc   152
   Leu Leu Leu Ser Ser Arg Thr Ala Arg Ser Glu Glu Asp Arg Asp Gly
        20                  25                  30
55
   cta tgg gat gcc tgg ggc cca tgg agt gaa tgc tca cgc acg tgt ggg   200
   Leu Trp Asp Ala Trp Gly Pro Trp Ser Glu Cys Ser Arg Thr Cys Gly
    35                  40                  45                  50

   gaa gga ata cag gtt gt  aa tta tgg cta tgt cgg cct gat gtt aca ca
   Glu Gly Arg Arg Ile Arg Tyr Arg Thr Cys Asn Val Arg Cys Pro
                   55                  60                  65
```

```
   cca gaa gca ggt gat ttc cga gct cag caa tgc tca gct cat aat gat   344
   Pro Glu Ala Gly Asp Phe Arg Ala Gln Gln Cys Ser Ala His Asn Asp
             85                  90                  95
5
   gtc aag cac cat ggc cag ttt tat gaa tgg ctt cct gtg tct aat gac   392
   Val Lys His His Gly Gln Phe Tyr Glu Trp Leu Pro Val Ser Asn Asp
         100                 105                 110

10 cct gac aac cca tgt tca ctc aag tgc caa gcc aaa gga aca acc ctg   440
   Pro Asp Asn Pro Cys Ser Leu Lys Cys Gln Ala Lys Gly Thr Thr Leu
   115                 120                 125                 130

   gtt gtt gaa cta gca cct aag gtc tta gat ggt acg cgt tgc tat aca   488
15 Val Val Glu Leu Ala Pro Lys Val Leu Asp Gly Thr Arg Cys Tyr Thr
                   135                 140                 145

   gaa tct ttg gat atg tgc atc agt ggt tta tgc caa att gtt ggc tgc   536
   Glu Ser Leu Asp Met Cys Ile Ser Gly Leu Cys Gln Ile Val Gly Cys
20             150                 155                 160

   gat cac cag ctg gga agc acc gtc aag gaa gat aac tgt ggg gtc tgc   584
   Asp His Gln Leu Gly Ser Thr Val Lys Glu Asp Asn Cys Gly Val Cys
           165                 170                 175
25
   aac gga gat ggg tcc acc tgc cgg ctg gtc cga ggg cag tat aaa tcc   632
   Asn Gly Asp Gly Ser Thr Cys Arg Leu Val Arg Gly Gln Tyr Lys Ser
         180                 185                 190

30 cag ctc tcc gca acc aaa tcg gat gat act gtg gtt gca att ccc tat   680
   Gln Leu Ser Ala Thr Lys Ser Asp Asp Thr Val Val Ala Ile Pro Tyr
   195                 200                 205                 210

   gga agt aga cat att cgc ctt gtc tta aaa ggt cct gat cac tta tat   728
35 Gly Ser Arg His Ile Arg Leu Val Leu Lys Gly Pro Asp His Leu Tyr
                   215                 220                 225

   ctg gaa acc aaa acc ctc cag ggg act aaa ggt gaa aac agt ctc agc   776
   Leu Glu Thr Lys Thr Leu Gln Gly Thr Lys Gly Glu Asn Ser Leu Ser
40             230                 235                 240

   tcc aca gga act ttc ctt gtg gac aat tct agt gtg gac ttc cag aaa   824
   Ser Thr Gly Thr Phe Leu Val Asp Asn Ser Ser Val Asp Phe Gln Lys
           245                 250                 255
45
   ttt cca gac aaa gag ata ctg aga atg gct gga ccg ttc aca gca gat   872
   Phe Pro Asp Lys Glu Ile Leu Arg Met Ala Gly Pro Leu Thr Ala Asp
         260                 265                 270

50 ttc att gtc aag att cgt aac tcg ggc ttc gct gac agt tca gtc cag   920
   Phe Ile Val Lys Ile Arg Asn Ser Gly Ser Ala Asp Ser Thr Val Gln
   275                 280                 285                 290

   ttc atc ttc tat caa ccc atc atc cac cga tgg agg gag acg gat ttc   968
55 Phe Ile Phe Tyr Gln Pro Ile Ile His Arg Trp Arg Glu Thr Asp Phe
                   295                 300                 305

   ... ... ... ... ... ... ... ... ... ... ... ... ... ... ...  1016

   ...                 ...                 ...
...
   ... ... ... ... ... ... ... ... ... ... ... ... ... ... ...  1112
```

```
   His Tyr Tyr Pro Glu Asn Ile Lys Pro Lys Pro Lys Leu Gln Glu Cys
       340                 345                 350

   aac ttg gat cct tgt cca gcc agt gac gga tac aag cag atc atg cct   1140
 5 Asn Leu Asp Pro Cys Pro Ala Ser Asp Gly Tyr Lys Gln Ile Met Pro
   355                 360                 365                 370

   tat gac ctc tac cat ccc ctt cct cgg tgg gag gcc acc cca tgg acc   1200
   Tyr Asp Leu Tyr His Pro Leu Pro Arg Trp Glu Ala Thr Pro Trp Thr
10                 375                 380                 385

   gcg tgc tcc tcc tcg tgt ggg ggg ggc atc cag agc cgg gca gtt tcc   1256
   Ala Cys Ser Ser Ser Cys Gly Gly Gly Ile Gln Ser Arg Ala Val Ser
               390                 395                 400
15
   tgt gtg gag gag gac atc cag ggg cat gtc act tca gtg gaa gag tgg   1304
   Cys Val Glu Glu Asp Ile Gln Gly His Val Thr Ser Val Glu Glu Trp
           405                 410                 415

20 aaa tgc atg tac acc cct aag atg ccc atc gcg cag ccc tgc aac att   1352
   Lys Cys Met Tyr Thr Pro Lys Met Pro Ile Ala Gln Pro Cys Asn Ile
       420                 425                 430

   ttt gac tgc cct aaa tgg ctg gca cag gag tgg tct ccg tgc aca gtg   1400
25 Phe Asp Cys Pro Lys Trp Leu Ala Gln Glu Trp Ser Pro Cys Thr Val
   435                 440                 445                 450

   acg tgt ggc cag ggc ctc aga tac cgt gtg gtc ctc tgc atc gac cat   1448
   Thr Cys Gly Gln Gly Leu Arg Tyr Arg Val Val Leu Cys Ile Asp His
30                 455                 460                 465

   cga gga atg cac aca gga ggc tgt agc cca aaa aca aag ccc cac ata   1496
   Arg Gly Met His Thr Gly Gly Cys Ser Pro Lys Thr Lys Pro His Ile
               470                 475                 480
35
   aaa gag gaa tgc atc gta ccc act ccc tgc tat aaa ccc aaa gag aaa   1544
   Lys Glu Glu Cys Ile Val Pro Thr Pro Cys Tyr Lys Pro Lys Glu Lys
           485                 490                 495

40 ctt cca gtc gag gcc aag ttg cca tgg ttc aaa caa gct caa gag cta   1592
   Leu Pro Val Glu Ala Lys Leu Pro Trp Phe Lys Gln Ala Gln Glu Leu
       500                 505                 510

   gaa gaa gga gct gct gtg tca gag gag ccc tcg taa gtt gta           1634
45 Glu Glu Gly Ala Ala Val Ser Glu Glu Pro Ser     Val Val
   515                 520                 525

   aaaggaaaga ctgttctatt tttgaaactt ttgttttaag aaaagcagtgt ctcactggtt 1694

91 gtagctttca tgggttctga actaagtgta atcatctcat taaagctttt tggctctaaa 1754

   atttaaagatt gatttagtttc aaaaaaaaaa aaaaaaaaga tgcggccgc           1803


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                                                                    .
.. Asp Gly Leu Trp Asp Val Trp Ile Glu Arg Ser Val Ile Arg Ile
          35                  40                  45
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Cys Gly Gly Gly Ala Ala Asn Ser Leu Arg Arg Cys Leu Ser Ser Lys
        50                  55                  60
Ser Cys Glu Gly Arg Asn Ile Arg Tyr Arg Thr Cys Ser Asn Val Asp
 65                  70                  75                  80
Cys Pro Pro Glu Ala Gly Asp Phe Arg Ala Gln Gln Cys Ser Ala His
                85                  90                  95
Asn Asp Val Lys His His Gly Gln Phe Tyr Glu Trp Leu Pro Val Ser
            100                 105                 110
Asn Asp Pro Asp Asn Pro Cys Ser Leu Lys Cys Gln Ala Lys Gly Thr
        115                 120                 125
Thr Leu Val Val Glu Leu Ala Pro Lys Val Leu Asp Gly Thr Arg Cys
    130                 135                 140
Tyr Thr Glu Ser Leu Asp Met Cys Ile Ser Gly Leu Cys Gln Ile Val
145                 150                 155                 160
Gly Cys Asp His Gln Leu Gly Ser Thr Val Lys Glu Asp Asn Cys Gly
                165                 170                 175
Val Cys Asn Gly Asp Gly Ser Thr Cys Arg Leu Val Arg Gly Gln Tyr
            180                 185                 190
Lys Ser Gln Leu Ser Ala Thr Lys Ser Asp Asp Thr Val Val Ala Ile
        195                 200                 205
Pro Tyr Gly Ser Arg His Ile Arg Leu Val Leu Lys Gly Pro Asp His
    210                 215                 220
Leu Tyr Leu Glu Thr Lys Thr Leu Gln Gly Thr Lys Gly Glu Asn Ser
225                 230                 235                 240
Leu Ser Ser Thr Gly Thr Phe Leu Val Asp Asn Ser Ser Val Asp Phe
                245                 250                 255
Gln Lys Phe Pro Asp Lys Glu Ile Leu Arg Met Ala Gly Pro Leu Thr
            260                 265                 270
Ala Asp Phe Ile Val Lys Ile Arg Asn Ser Gly Ser Ala Asp Ser Thr
        275                 280                 285
Val Gln Phe Ile Phe Tyr Gln Pro Ile Ile His Arg Trp Arg Glu Thr
    290                 295                 300
Asp Phe Phe Pro Cys Ser Ala Thr Cys Gly Gly Gly Tyr Gln Leu Thr
305                 310                 315                 320
Ser Ala Glu Cys Tyr Asp Leu Arg Ser Asn Arg Val Val Ala Asp Gln
                325                 330                 335
Tyr Cys His Tyr Tyr Pro Glu Asn Ile Lys Pro Lys Pro Lys Leu Gln
            340                 345                 350
Glu Cys Asn Leu Asp Pro Cys Pro Ala Ser Asp Gly Tyr Lys Gln Ile
        355                 360                 365
Met Pro Tyr Asp Leu Tyr His Pro Leu Pro Arg Trp Glu Ala Thr Pro
    370                 375                 380
Trp Thr Ala Cys Ser Ser Ser Cys Gly Gly Gly Ile Gln Ser Arg Ala
385                 390                 395                 400
Val Ser Cys Val Glu Glu Asp Ile Gln Gly His Val Thr Ser Val Glu
                405                 410                 415
Glu Trp Lys Cys Met Tyr Thr Pro Lys Met Pro Ile Ala Gln Pro Cys
            420                 425                 430
Asn Ile Phe Asp Cys Pro Lys Trp Leu Ala Gln Glu Trp Ser Pro Cys
        435                 440                 445
Thr Val Thr Cys Gly Gln Gly Leu Arg Tyr Arg Val Val Leu Cys Ile
    450                 455                 460
Asp His Arg Gly Met His Thr Gly Gly Cys Ser Pro Lys Thr Lys Pro
465                 470                 475                 480
His Ile Lys Glu Glu Cys Ile Val Pro Thr Pro Cys Tyr Lys Pro Lys
                485                 490                 495
Glu Lys Leu Pro Val Glu Ala Lys Leu Pro Trp Phe Lys Gln Ala Gln
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<213> Homo sapiens ADAMTS-5

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 1               5                  10                  15

Thr Phe Gly Ser Thr Glu Asp Lys Arg Leu Met Ser Ser Ile Leu Thr
            20                  25                  30

Ser Ile Asp Ala Ser Lys Pro Trp Ser Lys Cys Thr Ser Ala Thr Ile
        35                  40                  45

Thr Glu Phe Leu Asp Asp Gly His Gly Asn Cys Leu Leu Asp Leu Pro
    50                  55                  60

Arg Lys Gln Ile Leu Gly Pro Glu Glu Leu Pro Gly Gln Thr Tyr Asp
65                  70                  75                  80

Ala Thr Gln Gln Cys Asn Leu Thr Phe Gly Pro Asp Tyr Ser Val Cys
                85                  90                  95

Pro Gly Xaa Asp Val Cys Ala Arg Leu Trp Cys Ala Val Val Arg Gln
            100                 105                 110

Gly Gln Met Val Cys Leu Thr Lys Lys Leu Pro Ala Val Glu Gly Thr
        115                 120                 125

Pro Cys Gly Lys Gly Arg Ile Cys Leu Gln Gly Lys Cys Val Asp Lys
    130                 135                 140

Thr Lys Lys Lys Tyr Tyr Ser Thr Ser Ser His Gly Asn Trp Gly Ser
145                 150                 155                 160

Trp Gly Ser Trp Gly Gln Cys Ser Arg Ser Cys Gly Gly Gly Val Gln
                165                 170                 175

Phe Ala Tyr Arg His Cys Asn Asn Pro Ala Pro Arg Asn Asn Gly Arg
            180                 185                 190

Tyr Cys Thr Gly Lys Arg Ala Ile Tyr His Ser Cys Ser Leu Met Pro
        195                 200                 205

Cys Pro Pro Asn Gly Lys Ser Phe Arg His Glu Gln Cys Glu Ala Lys
    210                 215                 220

Asn Gly Tyr Gln Ser Asp Ala Lys Gly Val Lys Thr Phe Val Glu Trp
225                 230                 235                 240

Val Pro Lys Tyr Ala Gly Val Leu Pro Ala Asp Val Cys Lys Leu Thr
                245                 250                 255

Cys Arg Ala Lys Gly Thr Gly Tyr Tyr Val Val Phe Ser Pro Lys Val
            260                 265                 270

Thr Asp Gly Thr Glu Cys Arg Pro Tyr Ser Asn Ser Val Cys Val Arg
        275                 280                 285
```

Val Arg Ile Pro Glu Gly Ala Thr His Ile Lys Val Arg Gln Phe Lys
340 345 350

Ala Lys Asp Gln Thr Arg Phe Thr Ala Tyr Leu Ala Leu Lys Lys Lys
355 360 365

Asn Gly Glu Tyr Leu Ile Asn Gly Lys Tyr Met Ile Ser Thr Ser Glu
370 375 380

Thr Ile Ile Asp Ile Asn Gly Thr Val Met Asn Tyr Ser Gly Trp Ser
385 390 395 400

His Arg Asp Asp Phe Leu His Gly Met Gly Tyr Ser Ala Thr Lys Glu
405 410 415

Ile Leu Ile Val Gln Ile Leu Ala Thr Asp Pro Thr Lys Pro Leu Asp
420 425 430

Val Arg Tyr Ser Phe Phe Val Pro Lys Lys Ser Thr Pro Lys Val Asn
435 440 445

Ser Val Thr Ser His Gly Ser Asn Lys Val Gly Ser His Thr Ser Gln
450 455 460

Pro Gln Trp Val Thr Gly Pro Trp Leu Ala Cys Ser Arg Thr Cys Asp
465 470 475 480

Thr Gly Trp His Thr Arg Thr Val Gln Cys Gln Asp Gly Asn Arg Lys
485 490 495

Leu Ala Lys Gly Cys Pro Leu Ser Gln Arg Pro Ser Ala Phe Lys Gln
500 505 510

Cys Leu Leu Lys Lys Cys
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1 5

atg aga tcg cct tcc tca ccc gcg tgg acc aca acg ggg cac tgc tgg 99
Met Arg Ser Pro Ser Pro Pro Ala Trp Thr Thr Thr Gly His Cys Trp
10 15 20 25

cct tct cgc cac ctc ctc cct gga gca gcg ccg cgg cac ggg ggc cac 147
Pro Ser Arg His Leu Leu Pro Gly Ala Ala Pro Arg His Gly Gly His
30 35 40

Leu Leu Asp Leu Thr Arg Leu Tyr Arg Leu Ala Gly Arg Val
45 50

```
   gtg gag tac tgg aca cgg gag ggc ctg gcc tgg cag agg gcg gcc cgg   291
   Val Glu Tyr Trp Thr Arg Glu Gly Leu Ala Trp Gln Arg Ala Ala Arg
        75                  80                  85
 5
   ccc cac tgc ctc tac gct ggt cac ctg cag ggc cag gcc agc agc tcc   339
   Pro His Cys Leu Tyr Ala Gly His Leu Gln Gly Gln Ala Ser Ser Ser
    90                  95                 100                 105

10 cat gtg gcc atc agc acc tgt gga ggc ctg cac ggc ctg atc gtg gca   387
   His Val Ala Ile Ser Thr Cys Gly Gly Leu His Gly Leu Ile Val Ala
                   110                 115                 120

   gac gag gaa gag tac ctg att gag ccc ctg cac ggt ggg ccc aag ggt   435
15 Asp Glu Glu Glu Tyr Leu Ile Glu Pro Leu His Gly Gly Pro Lys Gly
               125                 130                 135

   tct cgg agc ccg gag gaa agt gga cca cat gtg gtg tac aag cgt tcc   483
   Ser Arg Ser Pro Glu Glu Ser Gly Pro His Val Val Tyr Lys Arg Ser
20         140                 145                 150

   tct ctg cgt cac ccc cac ctg gac aca gcc tgt gga gtg aga gat gag   531
   Ser Leu Arg His Pro His Leu Asp Thr Ala Cys Gly Val Arg Asp Glu
       155                 160                 165
25
   aaa ccg tgg aaa ggg cgg cca tgg tgg ctg cgg acc ttg aag cca ccg   579
   Lys Pro Trp Lys Gly Arg Pro Trp Trp Leu Arg Thr Leu Lys Pro Pro
   170                 175                 180                 185

30 cct gcc aga ccc ctg ggg aat gaa aca gag cgt ggt cag cca ggc ctg   627
   Pro Ala Arg Pro Leu Gly Asn Glu Thr Glu Arg Gly Gln Pro Gly Leu
                   190                 195                 200

   aag cga tcg gtc agc cga gag cgc tac gtg gag acc ctg gtg gtg gct   675
35 Lys Arg Ser Val Ser Arg Glu Arg Tyr Val Glu Thr Leu Val Val Ala
               205                 210                 215

   gac aag atg atg gtg gcc tat cac ggg cgc cgg gat gtg gag cag tat   723
   Asp Lys Met Met Val Ala Tyr His Gly Arg Arg Asp Val Glu Gln Tyr
40         220                 225                 230

   gtc ctg gcc atc atg aac att gtt gcc aaa ctt ttc cag gac tcg agt   771
   Val Leu Ala Ile Met Asn Ile Val Ala Lys Leu Phe Gln Asp Ser Ser
       235                 240                 245
45
   ctg gga agc acc gtt aac atc ctc gta act cgc ctt atc ctg ctt acg   819
   Leu Gly Ser Thr Val Asn Ile Leu Val Thr Arg Leu Ile Leu Leu Thr
   250                 255                 260                 265

50 gag gac cag ccg act ctg gag atc acc cac cat gcc ggg aag tcc ctg   867
   Glu Asp Gln Pro Thr Leu Glu Ile Thr His His Ala Gly Lys Ser Leu
                   270                 275                 280

   gac agc ttc tgt aag tgg cag aaa tcc atc gtg aac cac agc ggc cat   915
55 Asp Ser Phe Cys Lys Trp Gln Lys Ser Ile Val Asn His Ser Gly His
               285                 290                 295

   ggc ctc acc gag ctc ccc ttc gcc gag gcc ggt ctg acc gtg ggg gac   963

   ...                 ...                 ...
65
   acg ctg ggc atg gcc ggg tgt ggg gac ttt gtt gtg aac acg agc cac   ....
```

```
    Thr Leu Gly Leu Ala Arg Trp Ala Glu Cys Val Ser Ala Arg Glu Ala
    330                 335                 340                 345

    gca ggg tca atg agg aca ttg gct gcc aca agc gtt cac cat tgc cac   1107
 5  Ala Ala Ser Met Arg Thr Leu Ala Ala Thr Ser Val His His Cys His
                    350                 355                 360

    gag atc ggg cac aca ttc ggc atg aac cat gac ggc gtg gga aac agc   1155
    Glu Ile Gly His Thr Phe Gly Met Asn His Asp Gly Val Gly Asn Ser
10              365                 370                 375

    tgt ggg gcc cgt ggt cag gac cca gcc aag ctc atg gct gcc cac att   1203
    Cys Gly Ala Arg Gly Gln Asp Pro Ala Lys Leu Met Ala Ala His Ile
            380                 385                 390
15
    acc atg aag acc aac cca ttc gtg tgg tca tcc tgc aac cgt gac tac   1251
    Thr Met Lys Thr Asn Pro Phe Val Trp Ser Ser Cys Asn Arg Asp Tyr
        395                 400                 405

20  atc acc agc ttt cta gac tcg ggc ctg ggg ctc tgc ctg aac aac cgg   1299
    Ile Thr Ser Phe Leu Asp Ser Gly Leu Gly Leu Cys Leu Asn Asn Arg
    410                 415                 420                 425

    ccc ccc aga cag gac ttt gtg tac ccg aca gtg gca ccg ggc caa gcc   1347
25  Pro Pro Arg Gln Asp Phe Val Tyr Pro Thr Val Ala Pro Gly Gln Ala
                    430                 435                 440

    tac gat gca gat gag caa tgc cgc ttt cag cat gga gtc aaa tcg cgt   1395
    Tyr Asp Ala Asp Glu Gln Cys Arg Phe Gln His Gly Val Lys Ser Arg
30              445                 450                 455

    cag tgt aaa tac ggg gag gtc tgc agc gag ctg tgg tgt ctg agc aag   1443
    Gln Cys Lys Tyr Gly Glu Val Cys Ser Glu Leu Trp Cys Leu Ser Lys
            460                 465                 470
35
    agc aac cgg tgc atc acc aac agc atc ccg gcc gcc gag ggc acg ctg   1491
    Ser Asn Arg Cys Ile Thr Asn Ser Ile Pro Ala Ala Glu Gly Thr Leu
        475                 480                 485

40  tgc cag acg cac acc atc gac aag ggg tgg tgc tac aaa cgg gtc tgt   1539
    Cys Gln Thr His Thr Ile Asp Lys Gly Trp Cys Tyr Lys Arg Val Cys
    490                 495                 500                 505

    gtc ccc ttt ggg tcg cgc cca gag ggt gtg gac gga gcc tgg ggg ccg   1587
45  Val Pro Phe Gly Ser Arg Pro Glu Gly Val Asp Gly Ala Trp Gly Pro
                    510                 515                 520

    tgg act cca tgg ggc gac tgc agc cgg acc tgt ggc ggc ggc gtg tcc   1635
    Trp Thr Pro Trp Gly Asp Cys Ser Arg Thr Cys Gly Gly Gly Val Ser
50              525                 530                 535

    tct tct agt cgt cac tgc gac agc ccc agg cca acc atc ggg ggc aag   1683
    Ser Ser Ser Arg His Cys Asp Ser Pro Arg Pro Thr Ile Gly Gly Lys
            540                 545                 550
55
    tac tgt ctg ggt gag cga agg cgg cac cgc tcc tgc aac acg gat gac   1731
    Tyr Cys Leu Gly Glu Arg Arg Arg His Arg Ser Cys Asn Thr Asp Asp
        555                 560                 565

    gac agt ttc  tcc  ttc  gtg ggg aag gtg tac aag tgg tcc tcc acg gtg   1779
60  Asp Ser Phe  Ser  Phe  Val Gly Lys Val Tyr Lys Trp Ser Ser Thr Val
    570                 575                 580                 585
```

```
gga ggg ggc gtg aag gcc tgc tcg ctc acg agc cta gcg gaa ggc ttc   1875
Gly Gly Gly Val Lys Ala Cys Ser Leu Thr Ser Leu Ala Glu Gly Phe
            605                 610                 615

aac ttc tac acg gag agg gcg gca gcc gtg gtg gac ggg aca ccc tgc   1923
Asn Phe Tyr Thr Glu Arg Ala Ala Ala Val Val Asp Gly Thr Pro Cys
        620                 625                 630

cgt cca gac acg gtg gac att tgc gtc agt ggc gaa tgc aag cac gtg   1971
Arg Pro Asp Thr Val Asp Ile Cys Val Ser Gly Glu Cys Lys His Val
    635                 640                 645

ggc tgc gac cga gtc ctg ggc tcc gac ctg cgg gag gac aag tgc cga   2019
Gly Cys Asp Arg Val Leu Gly Ser Asp Leu Arg Glu Asp Lys Cys Arg
650                 655                 660                 665

gtg tgt ggc ggt gac ggc agt gcc tgc gag acc atc gag ggc gtc ttc   2067
Val Cys Gly Gly Asp Gly Ser Ala Cys Glu Thr Ile Glu Gly Val Phe
                670                 675                 680

agc cca gcc tca cct ggg gcc ggg tac gag gat gtc gtc tgg att ccc   2115
Ser Pro Ala Ser Pro Gly Ala Gly Tyr Glu Asp Val Val Trp Ile Pro
            685                 690                 695

aaa ggc tcc gtc cac atc ttc atc cag gat ctg aac ctc tct ctc agt   2163
Lys Gly Ser Val His Ile Phe Ile Gln Asp Leu Asn Leu Ser Leu Ser
        700                 705                 710

cac ttg gcc ctg aag gga gac cag gag tcc ctg ctg ctg gag ggg ctg   2211
His Leu Ala Leu Lys Gly Asp Gln Glu Ser Leu Leu Leu Glu Gly Leu
    715                 720                 725

cct ggg acc ccc cag ccc cac cgt ctg cct cta gct ggg acc acc ttt   2259
Pro Gly Thr Pro Gln Pro His Arg Leu Pro Leu Ala Gly Thr Thr Phe
730                 735                 740                 745

caa ctg cga cag ggg cca gac cag gtc cag agc ctc gaa gcc ctg gga   2307
Gln Leu Arg Gln Gly Pro Asp Gln Val Gln Ser Leu Glu Ala Leu Gly
                750                 755                 760

ccg att aat gca tct ctc atc gtc atg gtg ctg gcc cgg acc gag ctg   2355
Pro Ile Asn Ala Ser Leu Ile Val Met Val Leu Ala Arg Thr Glu Leu
            765                 770                 775

cct gcc ctc cgc tac cgc ttc aat gcc ccc atc gcc cgt gac tcg ctg   2403
Pro Ala Leu Arg Tyr Arg Phe Asn Ala Pro Ile Ala Arg Asp Ser Leu
        780                 785                 790

ccc ccc tac tcc tgg cac tat gcg ccc tgg acc aag tgc tcg gcc cag   2451
Pro Pro Tyr Ser Trp His Tyr Ala Pro Trp Thr Lys Cys Ser Ala Gln
    795                 800                 805

tgt gca ggc ggt agc cag gtg cag gcg gtg gag tgc cgc aac cag ctg   2499
Cys Ala Gly Gly Ser Gln Val Gln Ala Val Glu Cys Arg Asn Gln Leu
810                 815                 820                 825

[illegible]
```

```
   Val Val Gly Asn Trp Ser Leu Cys Ser Arg Ser Cys Asp Ala Gly Val
           860                 865                 870

   cgc agt cgc tcg gtc gtg tgc cag cgc cgc gtc tct gcc gcg gag gag   2691
 5 Arg Ser Arg Ser Val Val Cys Gln Arg Arg Val Ser Ala Ala Glu Glu
       875                 880                 885

   aag gcg ctg gac gac agc gca tgc ccg cag ccg cgc cca cct gta ctg   2739
   Lys Ala Leu Asp Asp Ser Ala Cys Pro Gln Pro Arg Pro Pro Val Leu
10 890                 895                 900                 905

   gag gcc tgc cac ggc ccc act tgc cct ccg gag tgg gcg gcc ctc gac   2787
   Glu Ala Cys His Gly Pro Thr Cys Pro Pro Glu Trp Ala Ala Leu Asp
                   910                 915                 920
15
   tgg tct gag tgc acc ccc agc tgc ggg ccg ggc ctc cgc cac cgc gtg   2835
   Trp Ser Glu Cys Thr Pro Ser Cys Gly Pro Gly Leu Arg His Arg Val
               925                 930                 935

20 gtc ctt tgc aag agc gca gac cac cgc gcc acg ctg ccc ccg gcg cac   2883
   Val Leu Cys Lys Ser Ala Asp His Arg Ala Thr Leu Pro Pro Ala His
           940                 945                 950

   tgc tca ccc gcc gcc aag cca ccg gct acc atg cgc tgc aac ttg cgc   2931
25 Cys Ser Pro Ala Ala Lys Pro Pro Ala Thr Met Arg Cys Asn Leu Arg
       955                 960                 965

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30 970                 975                 980                 985

   gca cag tgc ggc gtc ggg cag cgg cag cgc tcg gtg cgc tgc acc agc   3027
   Ala Gln Cys Gly Val Gly Gln Arg Gln Arg Ser Val Arg Cys Thr Ser
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   cac acg ggc cag gcg tcg cac gag tgc acg gag gcc ctg cgg ccg ccc   3075
   His Thr Gly Gln Ala Ser His Glu Cys Thr Glu Ala Leu Arg Pro Pro
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40 acc acg cag cag tgt gag gcc aag tgc gac agc cca acc ccc ggg gac   3123
   Thr Thr Gln Gln Cys Glu Ala Lys Cys Asp Ser Pro Thr Pro Gly Asp
           1020                1025                1030

   ggc cct gaa gag tgc aag gat gtg aac aag gtc gcc tac tgc ccc ctg   3171
45 Gly Pro Glu Glu Cys Lys Asp Val Asn Lys Val Ala Tyr Cys Pro Leu
       1035                1040                1045

   gtg ctc aaa ttt cag ttc tgc agc cga gcc tac ttc cgc cag atg tgc   3219
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50 1050                1055                1060                1065

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   Cys Lys Thr Cys Gln Gly His
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Gln Ser Gly Leu Ala Ser Thr His Phe Leu Leu Asn Leu Thr Arg Ser
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Ser Arg Leu Leu Ala Gly Arg Val Ser Val Glu Tyr Trp Thr Arg Glu
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Gly Leu Ala Trp Gln Arg Ala Ala Arg Pro His Cys Leu Tyr Ala Gly
                 85                  90                  95

His Leu Gln Gly Gln Ala Ser Ser Ser His Val Ala Ile Ser Thr Cys
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Gly Gly Leu His Gly Leu Ile Val Ala Asp Glu Glu Glu Tyr Leu Ile
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Glu Pro Leu His Gly Gly Pro Lys Gly Ser Arg Ser Pro Glu Glu Ser
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Gly Pro His Val Val Tyr Lys Arg Ser Ser Leu Arg His Pro His Leu
145                 150                 155                 160

Asp Thr Ala Cys Gly Val Arg Asp Glu Lys Pro Trp Lys Gly Arg Pro
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Trp Trp Leu Arg Thr Leu Lys Pro Pro Pro Ala Arg Pro Leu Gly Asn
            180                 185                 190

Glu Thr Glu Arg Gly Gln Pro Gly Leu Lys Arg Ser Val Ser Arg Glu
        195                 200                 205

Arg Tyr Val Glu Thr Leu Val Val Ala Asp Lys Met Met Val Ala Tyr
    210                 215                 220

His Gly Arg Arg Asp Val Glu Gln Tyr Val Leu Ala Ile Met Asn Ile
225                 230                 235                 240

Val Ala Lys Leu Phe Gln Asp Ser Ser Leu Gly Ser Thr Val Asn Ile
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Leu Val Thr Arg Leu Ile Leu Leu Thr Glu Asp Gln Pro Thr Leu Glu
            260                 265                 270

Ile Thr His His Ala Gly Lys Ser Leu Asp Ser Phe Cys Lys Trp Gln
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Lys Ser Ile Val Asn His Ser Gly His Gly Asn Ala Ile Pro Glu Asn
    290                 295                 300

Ala Glu Cys Val Ser Ala Arg Glu Ala Ala Ala Ser Met Arg Thr Leu
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                340                 345                 350

   Ala Ala Thr Ser Val His His Cys His Glu Ile Gly His Thr Phe Gly
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   Pro Ala Lys Leu Met Ala Ala His Ile Thr Met Lys Thr Asn Pro Phe
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   Val Trp Ser Ser Cys Asn Arg Asp Tyr Ile Thr Ser Phe Leu Asp Ser
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15 Gly Leu Gly Leu Cys Leu Asn Asn Arg Pro Pro Arg Gln Asp Phe Val
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   Tyr Pro Thr Val Ala Pro Gly Gln Ala Tyr Asp Ala Asp Glu Gln Cys
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   Arg Phe Gln His Gly Val Lys Ser Arg Gln Cys Lys Tyr Gly Glu Val
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   Cys Ser Glu Leu Trp Cys Leu Ser Lys Ser Asn Arg Cys Ile Thr Asn
25 465                 470                 475                 480

   Ser Ile Pro Ala Ala Glu Gly Thr Leu Cys Gln Thr His Thr Ile Asp
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30 Lys Gly Trp Cys Tyr Lys Arg Val Cys Val Pro Phe Gly Ser Arg Pro
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   Glu Gly Val Asp Gly Ala Trp Gly Pro Trp Thr Pro Trp Gly Asp Cys
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40 545                 550                 555                 560

   Arg His Arg Ser Cys Asn Thr Asp Asp Cys Pro Pro Gly Ser Gln Asp
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45 Phe Arg Glu Val Gln Cys Ser Glu Phe Asp Ser Ile Pro Phe Arg Gly
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   Lys Phe Tyr Lys Trp Lys Thr Tyr Arg Gly Gly Gly Val Lys Ala Cys
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50
   Ser Leu Thr Ser Leu Ala Glu Gly Phe Asn Phe Tyr Thr Glu Arg Ala
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   Ala Ala Val Val Asp Gly Thr Pro Cys Arg Pro Asp Thr Val Asp Ile
55 625                 630                 635                 640

   Cys Val Ser Gly Glu Cys Lys His Val Gly Cys Asp Arg Val Leu Gly
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   Gly Tyr Glu Asp Val Val Trp Ile Pro Lys Gly Ser Val His Ile Thr
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690 695 700

Ile Gln Asp Leu Asn Leu Ser Leu Ser His Leu Ala Leu Lys Gly Asp
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Gln Glu Ser Leu Leu Leu Glu Gly Leu Pro Gly Thr Pro Gln Pro His
725 730 735

Arg Leu Pro Leu Ala Gly Thr Thr Phe Gln Leu Arg Gln Gly Pro Asp
740 745 750

Gln Val Gln Ser Leu Glu Ala Leu Gly Pro Ile Asn Ala Ser Leu Ile
755 760 765

Val Met Val Leu Ala Arg Thr Glu Leu Pro Ala Leu Arg Tyr Arg Phe
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Asn Ala Pro Ile Ala Arg Asp Ser Leu Pro Pro Tyr Ser Trp His Tyr
785 790 795 800

Ala Pro Trp Thr Lys Cys Ser Ala Gln Cys Ala Gly Gly Ser Gln Val
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Gln Ala Val Glu Cys Arg Asn Gln Leu Asp Ser Ser Ala Val Ala Pro
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His Tyr Cys Ser Ala His Ser Lys Leu Pro Lys Arg Gln Arg Ala Cys
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Asn Thr Glu Pro Cys Pro Pro Asp Trp Val Val Gly Asn Trp Ser Leu
850 855 860

Cys Ser Arg Ser Cys Asp Ala Gly Val Arg Ser Arg Ser Val Val Cys
865 870 875 880

Gln Arg Arg Val Ser Ala Ala Glu Glu Lys Ala Leu Asp Asp Ser Ala
885 890 895

Cys Pro Gln Pro Arg Pro Pro Val Leu Glu Ala Cys His Gly Pro Thr
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Cys Pro Pro Glu Trp Ala Ala Leu Asp Trp Ser Glu Cys Thr Pro Ser
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Cys Gly Pro Gly Leu Arg His Arg Val Val Leu Cys Lys Ser Ala Asp
930 935 940

His Arg Ala Thr Leu Pro Pro Ala His Cys Ser Pro Ala Ala Lys Pro
945 950 955 960

Pro Ala Thr Met Arg Cys Asn Leu Arg Arg Cys Pro Pro Ala Arg Trp
965 970 975

Val Ala Gly Glu Trp Gly Glu Cys Ser Ala Gln Cys Gly Val Gly Gln
980 985 990

Arg Gln Arg Ser Val Arg Cys Thr Ser His Thr Gly Gln Ala Ser His
995 1000 1005

Val Asn Lys Thr Ala Tyr Cys Pro Leu Leu Val Leu Lys Pro Gln Ser Cy

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   Thr Leu Leu Thr Leu Leu Val Arg Asp Leu Ala Glu Met Gly Ser Pro
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   gac gcc gcg gcg gcc gtg cgc aag gac agg ctg cac ccg agg caa gtg   149
   Asp Ala Ala Ala Ala Val Arg Lys Asp Arg Leu His Pro Arg Gln Val
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30 aaa tta tta gag acc ctg agc gaa tac gaa atc gtg tct ccc atc cga   197
   Lys Leu Leu Glu Thr Leu Ser Glu Tyr Glu Ile Val Ser Pro Ile Arg
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   gtg aac gct ctc gga gaa ccc ttt ccc acg aac gtc cac ttc aaa aga   245
35 Val Asn Ala Leu Gly Glu Pro Phe Pro Thr Asn Val His Phe Lys Arg
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   acg cga cgg agc att aac tct gcc act gac ccc tgg cct gcc ttc gcc   293
   Thr Arg Arg Ser Ile Asn Ser Ala Thr Asp Pro Trp Pro Ala Phe Ala
40              75                  80                  85

   tcc tcc tct tcc tcc tct acc tcc ccc cag gcg cat tac cgc ctc tct   341
   Ser Ser Ser Ser Ser Ser Thr Ser Pro Gln Ala His Tyr Arg Leu Ser
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   gcc ttc ggc cag cag ttt cta ttt aat ctc acc gcc aat gct gga ttt   389
   Ala Phe Gly Gln Gln Phe Leu Phe Asn Leu Thr Ala Asn Ala Gly Phe
       105                 110                 115

50 atc gct cca ctg ttc act gtc acc ctc ctc ggg acg ccc ggg gtg aat   437
   Ile Ala Pro Leu Phe Thr Val Thr Leu Leu Gly Thr Pro Gly Val Asn
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   cag acc aag ttt tat tcc gaa gag gaa gcg gaa ctc aag cac tgt ttc   485
55 Gln Thr Lys Phe Tyr Ser Glu Glu Glu Ala Glu Leu Lys His Cys Phe
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                   170                 175                 180
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    Phe Ile Glu Pro Leu Gln Ser Met Asp Glu Gln Glu Asp Glu Glu Glu
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    caa aac aaa ccc cac atc att tat agg cgc agc gcc ccc cag aga gag   577
 5  Gln Asn Lys Pro His Ile Ile Tyr Arg Arg Ser Ala Pro Gln Arg Glu
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    ccc tca aca gga agg cat gca tgt gac acc tca gaa cac aaa aat agg   625
    Pro Ser Thr Gly Arg His Ala Cys Asp Thr Ser Glu His Lys Asn Arg
10                  220                 225                 230

    cac agt aaa gac aag aag aaa acc aga gca aga aaa tgg gga gaa agg   673
    His Ser Lys Asp Lys Lys Lys Thr Arg Ala Arg Lys Trp Gly Glu Arg
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    att aac ctg gct ggt gac gta gca gca tta aac agc ggc tta gca aca   721
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20  gag gca ttt tct gct tat ggt aat aag acg gac aac aca aga gaa aag   769
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    agg acc cac aga agg aca aaa cgt ttt tta tcc tat cca cgg ttt gta   817
25  Arg Thr His Arg Arg Thr Lys Arg Phe Leu Ser Tyr Pro Arg Phe Val
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    gaa gtc ttg gtg gtg gca gac aac aga atg gtt tca tac cat gga gaa   865
    Glu Val Leu Val Val Ala Asp Asn Arg Met Val Ser Tyr His Gly Glu
30                  300                 305                 310

    aac ctt caa cac tat att tta act tta atg tca att gta gcc tct atc   913
    Asn Leu Gln His Tyr Ile Leu Thr Leu Met Ser Ile Val Ala Ser Ile
                315                 320                 325
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    tat aaa gac cca agt att gga aat tta att aat att gtt att gtg aac   961
    Tyr Lys Asp Pro Ser Ile Gly Asn Leu Ile Asn Ile Val Ile Val Asn
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40  tta att gtg att cat aat gaa cag gat ggg cct tcc ata tct ttt aat  1009
    Leu Ile Val Ile His Asn Glu Gln Asp Gly Pro Ser Ile Ser Phe Asn
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    gct cag aca aca tta aaa aac ttt tgc cag tgg cag cat tcg aac agt  1057
45  Ala Gln Thr Thr Leu Lys Asn Phe Cys Gln Trp Gln His Ser Asn Ser
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    cca ggt gga att cat tat gat act gct gtt ctt tta aca aga cag gat  1105
    Pro Gly Gly Ile His His Asp Thr Ala Val Leu Leu Thr Arg Gln Asp
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    atc tgc aga gct cac gac aaa tgt gat acc tta ggc ctg gct gaa ctg  1153
    Ile Cys Arg Ala His Asp Lys Cys Asp Thr Leu Gly Leu Ala Glu Leu
                395                 400                 405
55
    gga acc att tgt gat ccc tat aga agc tgt tct att agt gaa gat agt  1201
    Gly Thr Ile Cys Asp Pro Tyr Arg Ser Cys Ser Ile Ser Glu Asp Ser
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    gtt atc ttt tat gat gat aat aaa tat aaa tta gaa gga gtt gtt aaa  1249
    Val Ile Phe Tyr Asp Asp Asn Lys Tyr Lys Leu Glu Gly Val Val Lys
    425                 430                 435                 440
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475 480 485

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Thr Gly Tyr Gly Glu Cys Leu Leu Asn Glu Pro Glu Ser Arg Pro Tyr
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cct ttg cct gtc caa ctg cca ggc atc ctt tac aac gtg aat aaa caa 1589
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Gln Cys Arg Arg Leu Trp Ser Asn Asn Val Asn Gly Val His Lys Gly
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Cys Arg Thr Gln His Thr Pro Trp Ala Asp Gly Thr Glu Cys Glu Pro
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585 590 595

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Ser Arg Thr Cys Gly Gly Gly Ile Lys Thr Ala Ile Arg Glu Cys Asn
600 605 610 615

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635 640 645

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 5 Cys Asp His Val Leu Asn Ser Lys Ala Arg Arg Asp Lys Cys Gly Val
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   tgt ggt ggc gat aat tct tca tgc aaa aca gtg gca gga aca ttt aat   2309
   Cys Gly Gly Asp Asn Ser Ser Cys Lys Thr Val Ala Gly Thr Phe Asn
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   aca gta cat tat ggt tac aat act gtg gtc cga att cca gct ggt gct   2357
   Thr Val His Tyr Gly Tyr Asn Thr Val Val Arg Ile Pro Ala Gly Ala
   760                 765                 770                 775
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   acc aat att gat gtg cgg cag cac agt ttc tca ggg gaa aca gac gat   2405
   Thr Asn Ile Asp Val Arg Gln His Ser Phe Ser Gly Glu Thr Asp Asp
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   Asp Asn Tyr Leu Ala Leu Ser Ser Ser Lys Gly Glu Phe Leu Leu Asn
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   gga aac ttt gtt gtc aca atg gcc aaa agg gaa att cgc att ggg aat   2501
25 Gly Asn Phe Val Val Thr Met Ala Lys Arg Glu Ile Arg Ile Gly Asn
          810                 815                 820

   gct gtg gta gag tac agt ggg tcc gag act gcc gta gaa aga att aac   2549
   Ala Val Val Glu Tyr Ser Gly Ser Glu Thr Ala Val Glu Arg Ile Asn
30     825                 830                 835

   tca aca gat cgc att gag caa gaa ctt ttg ctt cag gtt ttg tcg gtg   2597
   Ser Thr Asp Arg Ile Glu Gln Glu Leu Leu Leu Gln Val Leu Ser Val
   840                 845                 850                 855
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   gga aag ttg tac aac ccc gat gta cgc tat tct ttc aat att cca att   2645
   Gly Lys Leu Tyr Asn Pro Asp Val Arg Tyr Ser Phe Asn Ile Pro Ile
                  860                 865                 870

40 gaa gat aaa cct cag cag ttt tac tgg aac agt cat ggg cca tgg caa   2693
   Glu Asp Lys Pro Gln Gln Phe Tyr Trp Asn Ser His Gly Pro Trp Gln
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   gca tgc agt aaa ccc tgc caa ggg gaa cgg aaa cga aaa ctt gtt tgc   2741
45 Ala Cys Ser Lys Pro Cys Gln Gly Glu Arg Lys Arg Lys Leu Val Cys
          890                 895                 900

   acc agg gaa tct gat cag ctt act gtt tct gat caa aga tgc gat cgg   2789
   Thr Arg Glu Ser Asp Gln Leu Thr Val Ser Asp Gln Arg Cys Asp Arg
50     905                 910                 915

   ctg ccc cag cct gga cac att act gaa ccc tgt ggt aca ggc tgt gac   2837
   Leu Pro Gln Pro Gly His Ile Thr Glu Pro Cys Gly Thr Gly Cys Asp
   920                 925                 930                 935
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   gta agg tgg cat gtt gcc agc agg agt gaa tgt agt gcc cag tgt ggc   2885
   Leu Arg Trp His Val Ala Ser Arg Ser Glu Cys Ser Ala Gln Cys Gly
                  940                 945                 950

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   tgg cgc tat tct gcc tgg act gaa tgt tca aaa agc tgt gac ggt ggg   3077
   Trp Arg Tyr Ser Ala Trp Thr Glu Cys Ser Lys Ser Cys Asp Gly Gly
  1000                1005                1010                1015

10 acc cag agg aga agg gct att tgt gtc aat acc cga aat gat gta ctg   3125
   Thr Gln Arg Arg Arg Ala Ile Cys Val Asn Thr Arg Asn Asp Val Leu
                  1020                1025                1030

   gat gac agc aaa tgc aca cat caa gag aaa gtt acc att cag agg tgc   3173
15 Asp Asp Ser Lys Cys Thr His Gln Glu Lys Val Thr Ile Gln Arg Cys
              1035                1040                1045

   agt gag ttc cct tgt cca cag tgg aaa tct gga gac tgg tca gag tgc   3221
   Ser Glu Phe Pro Cys Pro Gln Trp Lys Ser Gly Asp Trp Ser Glu Cys
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   ttg gtc acc tgt gga aaa ggg cat aag cac cgc cag gtc tgg tgt cag   3269
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     1065                1070                1075
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30 cca aca tct atg cag act tgt cag cag ccg gaa atg gca tcc tgg cag   3365
   Pro Thr Ser Met Gln Thr Cys Gln Gln Pro Glu Met Ala Ser Trp Gln
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   gcg ggt ccc tgg gta cag tgc agt gtc act tgt gga cag gga tac cag   3413
35 Ala Gly Pro Trp Val Gln Cys Ser Val Thr Cys Gly Gln Gly Tyr Gln
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   Leu Arg Ala Val Lys Cys Ile Ile Gly Thr Tyr Met Ser Val Val Asp
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   gac aat gac tgt aat gca gca act aga cca act gat acc cag gac tgt   3509
   Asp Asn Asp Cys Asn Ala Ala Thr Arg Pro Thr Asp Thr Gln Asp Cys
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   gaa tta cca tca tgt cat cct ccc cca gct gcw ccg gaa acg agg aga   3557
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50 agc aca tac agt gca cca aga acc cag tgg cga ttt ggg tct tgg act   3605
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   ggt agg gca acc cgg caa gtg atg tgt gtc aac tac agt gac cac gtg   3845
 5 Gly Arg Ala Thr Arg Gln Val Met Cys Val Asn Tyr Ser Asp His Val
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   atc gat cgg agt gag tgt gac cag gat tat atc cca gaa act gac cag   3893
   Ile Asp Arg Ser Glu Cys Asp Gln Asp Tyr Ile Pro Glu Thr Asp Gln
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   gac tgt tcc atg tca cca tgc cct caa agg acc cca gac agt ggc tta   3941
   Asp Cys Ser Met Ser Pro Cys Pro Gln Arg Thr Pro Asp Ser Gly Leu
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   gct cag cac ccc ttc caa aat gag gac tat cgt ccc cgg agc gcc agc   3989
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      1305                1310                1315

20 ccc agc cgc acc cat gtg ctc ggt gga aac cag tgg aga act ggc ccc   4037
   Pro Ser Arg Thr His Val Leu Gly Gly Asn Gln Trp Arg Thr Gly Pro
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   tgg gga gca tgt tcc agt acc tgt ggt ggc gga tcc cag cgg cgt gtt   4085
25 Trp Gly Ala Cys Ser Ser Thr Cys Ala Gly Gly Ser Gln Arg Arg Val
                  1340                1345                1350

   gtt gta tgt cag gat gaa aat gga tac acc gca aac gac tgt gtg gag   4133
   Val Val Cys Gln Asp Glu Asn Gly Tyr Thr Ala Asn Asp Cys Val Glu
30            1355                1360                1365

   aga ata aaa cct gat gag caa aga gcc tgt gaa tcc ggc cct tgt cct   4181
   Arg Ile Lys Pro Asp Glu Gln Arg Ala Cys Glu Ser Gly Pro Cys Pro
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   cag tgg gct tat ggc aac tgg gga gag tgc act aag ctg tgt ggt gga   4229
   Gln Trp Ala Tyr Gly Asn Trp Gly Glu Cys Thr Lys Leu Cys Gly Gly
      1385                1390                1395

40 ggc ata aga aca aga ctg gtg gtc tgt cag cgg tcc aac ggt gaa cgg   4277
   Gly Ile Arg Thr Arg Leu Val Val Cys Gln Arg Ser Asn Gly Glu Arg
   1400                1405                1410                1415

   ttt cca gat ttg agc tgt gaa att ctt gat aaa cct ccc gat cgt gag   4325
45 Phe Pro Asp Leu Ser Cys Glu Ile Leu Asp Lys Pro Pro Asp Arg Glu
                  1420                1425                1430

   cag tgt aac acg cat gct tgt cca cac gac gct gca tgg agt act ggc   4373
   Gln Cys Asn Thr His Ala Cys Pro His Asp Ala Ala Trp Ser Thr Gly
50            1435                1440                1445

   cct tgg agc tcg tgt tct gtc tct tgt ggt cga ggg cat aaa caa cga   4421
   Pro Trp Ser Ser Cys Ser Val Ser Cys Gly Arg Gly His Lys Gln Arg
         1450                1455                1460
55
   aat gtt tac tgc atg gca aaa gat gga agc cat tta gaa agt gat tac   4469
   Asn Val Tyr Cys Met Ala Lys Asp Gly Ser His Leu Glu Ser Asp Tyr

   [illegible]
   Arg Tyr Tyr Lys Trp Lys Ala Gly Ala Trp Val Ser Cys Ser Val Thr
                  1500                1505                1510
```

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   tgt ggc cga ggc gta cag cag agg cat gtg ggc tgt cag atc gga aca   4613
   Cys Gly Arg Gly Val Gln Gln Arg His Val Gly Cys Gln Ile Gly Thr
             1515                1520                1525
 5
   cac aaa ata gcc aga gac acc gag tgc aac cca tac acc aga ccg gag   4661
   His Lys Ile Ala Arg Asp Thr Glu Cys Asn Pro Tyr Thr Arg Pro Glu
          1530                1535                1540

10 tcg gaa tgc gaa tgc caa ggc cca cgg tgt ccc ctt tac act tgg agg   4709
   Ser Glu Cys Glu Cys Gln Gly Pro Arg Cys Pro Leu Tyr Thr Trp Arg
      1545                1550                1555

   gca gag gaa tcg caa gaa tgc acc aag acc tgc ggc gaa ggc tcc agg   4757
15 Ala Glu Glu Ser Gln Glu Cys Thr Lys Thr Cys Gly Glu Gly Ser Arg
   1560                1565                1570                1575

   tac cgc aaq gtq qtq tgt gtg gat gac aac aaa aac gag gtg cat ggg   4805
   Tyr Arg Lys Val Val Cys Val Asp Asp Asn Lys Asn Glu Val His Gly
20                 1580                1585                1590

   gca cgc tgt gac gtg agc aag cgg ccg gtg gac cgt gaa agc tgt agt   4853
   Ala Arg Cys Asp Val Ser Lys Arg Pro Val Asp Arg Glu Ser Cys Ser
             1595                1600                1605
25
   ttg caa ccc tgc gag tat gtc tgg atc aca gga gaa tgg tca gag tgc   4901
   Leu Gln Pro Cys Glu Tyr Val Trp Ile Thr Gly Glu Trp Ser Glu Cys
          1610                1615                1620

30 tca gtg acc tgt gga aaa ggc tac aaa caa agg ctt gtc tcg tgc agc   4949
   Ser Val Thr Cys Gly Lys Gly Tyr Lys Gln Arg Leu Val Ser Cys Ser
      1625                1630                1635

   gag att tac acc ggg aaa gag aat tat gaa tac agc tac caa acc acc   4997
35 Glu Ile Tyr Thr Gly Lys Glu Asn Tyr Glu Tyr Ser Tyr Gln Thr Thr
   1640                1645                1650                1655

   atc aac tgc cca ggc acg cag ccc ccc agt gtt cac ccc tgt tac ctg   5045
   Ile Asn Cys Pro Gly Thr Gln Pro Pro Ser Val His Pro Cys Tyr Leu
40                 1660                1665                1670

   agg gag tgc cct gtc tcg gcc acc tgg aga gtt ggc aac tgg ggg agc   5093
   Arg Glu Cys Pro Val Ser Ala Thr Trp Arg Val Gly Asn Trp Gly Ser
             1675                1680                1685
45
   tgc tca gtg tct tgt ggt gtt gga gtg atg cag aga tct gtg caa tgt   5141
   Cys Ser Val Ser Cys Gly Val Gly Val Met Gln Arg Ser Val Gln Cys
          1690                1695                1700

50 tta aac aat gag gac caa ccc agc cac tta tgc cac act gat ctg aag   5189
   Leu Thr Asn Glu Asp Gln Pro Ser His Leu Cys His Thr Asp Leu Lys
      1705                1710                1715

   cca gaa gaa cga aaa acc tgc cgt aat gtc tat aac tgt gag tta ccc   5237
55 Pro Glu Glu Arg Lys Thr Cys Arg Asn Val Tyr Asn Cys Glu Leu Pro
   1720                1725                1730                1735

   . . . . . .  . . . . . .  . . . . . .  . . . . . .  . . . . . .  .

   . . . . . .  . . . . . .  . . . . . .  . . . . . .  . . . . . .  .

   agg atg aac tgt gat cag ctg gaa gaa tca gtg aat gca   5281
```

```
   Gly Met His Ser Asp His Pro Lys Glu Tyr Val Thr Leu Val His Gly
          1770                1775                1780

   gac tct gag aat ttc tcc gag gtt tat ggg cac agg tta cac aac cca   5429
 5 Asp Ser Glu Asn Phe Ser Glu Val Tyr Gly His Arg Leu His Asn Pro
      1785                1790                1795

   aca gaa tgt ccc tat aac ggg agc cgg cgc gat gac tgc caa tgt cgg   5477
   Thr Glu Cys Pro Tyr Asn Gly Ser Arg Arg Asp Asp Cys Gln Cys Arg
10 1800                1805                1810                1815

   aag gat tac acg gcc gct ggg ttt tcc agt ttt cag aaa atc aga ata   5525
   Lys Asp Tyr Thr Ala Ala Gly Phe Ser Ser Phe Gln Lys Ile Arg Ile
                  1820                1825                1830
15
   gac ctg acc agc atg cag ata atc acc act gac tta cag ttt gca agg   5573
   Asp Leu Thr Ser Met Gln Ile Ile Thr Thr Asp Leu Gln Phe Ala Arg
             1835                1840                1845

20 aca agc gaa gga cat ccc gtc cct ttt gcc aca gcc ggg gat tgc tac   5621
   Thr Ser Glu Gly His Pro Val Pro Phe Ala Thr Ala Gly Asp Cys Tyr
          1850                1855                1860

   agc gct gcc aag tgc cca cag ggt cgt ttt agc atc aac ctt tat gga   5669
25 Ser Ala Ala Lys Cys Pro Gln Gly Arg Phe Ser Ile Asn Leu Tyr Gly
      1865                1870                1875

   acc ggc ttg tct tta act gaa tct gcc aga tgg ata tca caa ggg aat   5717
   Thr Gly Leu Ser Leu Thr Glu Ser Ala Arg Trp Ile Ser Gln Gly Asn
30 1880                1885                1890                1895

   tat gct gtc tct gac att aag aag tcg ccg gat ggt acc cga gtc gta   5765
   Tyr Ala Val Ser Asp Ile Lys Lys Ser Pro Asp Gly Thr Arg Val Val
                  1900                1905                1910
35
   ggg aaa tgc ggt ggt tac tgt gga aaa tgc act cca tcc tct ggt act   5813
   Gly Lys Cys Gly Gly Tyr Cys Gly Lys Cys Thr Pro Ser Ser Gly Thr
             1915                1920                1925

40 ggc ctg gag gtg cga gtt tta tagctaaggt gctttgaaga ggaagccatt      5864
   Gly Leu Glu Val Arg Val Leu
          1930

   atggatggat gaaggatagt aatgcaatac ctccaccttg atttgggtgc atgtgtatgt 5924
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   tatata                                                          5990

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   <213> Homo sapiens ADAMTS-9b
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   <400> 26
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    1               5                  10                  15

                    20                  25                  30

   Glu Ile Val Leu Pro Ile Arg Val Asn Ala Leu Gly Gln Pro Phe Pro
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50 55 60

Thr Asn Val His Phe Lys Arg Thr Arg Arg Ser Ile Asn Ser Ala Thr
65 70 75 80

Asp Pro Trp Pro Ala Phe Ala Ser Ser Ser Ser Ser Ser Thr Ser Pro
85 90 95

Gln Ala His Tyr Arg Leu Ser Ala Phe Gly Gln Gln Phe Leu Phe Asn
100 105 110

Leu Thr Ala Asn Ala Gly Phe Ile Ala Pro Leu Phe Thr Val Thr Leu
115 120 125

Leu Gly Thr Pro Gly Val Asn Gln Thr Lys Phe Tyr Ser Glu Glu Glu
130 135 140

Ala Glu Leu Lys His Cys Phe Tyr Lys Gly Tyr Val Asn Thr Asn Ser
145 150 155 160

Glu His Thr Ala Val Ile Ser Leu Cys Ser Gly Met Leu Gly Thr Phe
165 170 175

Arg Ser His Asp Gly Gly Tyr Phe Ile Glu Pro Leu Gln Ser Met Asp
180 185 190

Glu Gln Glu Asp Glu Glu Glu Gln Asn Lys Pro His Ile Ile Tyr Arg
195 200 205

Arg Ser Ala Pro Gln Arg Glu Pro Ser Thr Gly Arg His Ala Cys Asp
210 215 220

Thr Ser Glu His Lys Asn Arg His Ser Lys Asp Lys Lys Lys Thr Arg
225 230 235 240

Ala Arg Lys Trp Gly Glu Arg Ile Asn Leu Ala Gly Asp Val Ala Ala
245 250 255

Leu Asn Ser Gly Leu Ala Thr Glu Ala Phe Ser Ala Tyr Gly Asn Lys
260 265 270

Thr Asp Asn Thr Arg Glu Lys Arg Thr His Arg Arg Thr Lys Arg Phe
275 280 285

Leu Ser Tyr Pro Arg Phe Val Glu Val Leu Val Val Ala Asp Asn Arg
290 295 300

Met Val Ser Tyr His Gly Glu Asn Leu Gln His Tyr Ile Leu Thr Leu
305 310 315 320

Met Ser Ile Val Ala Ser Ile Tyr Lys Asp Pro Ser Ile Gly Asn Leu
325 330 335

Ile Asn Ile Val Ile Val Asn Leu Ile Val Ile His Asn Glu Gln Asp
340 345 350

Gly Pro Ser Ile Ser Phe Asn Ala Gln Thr Thr Leu Lys Asn Phe Cys
355 360 365

Thr Leu Gly Leu Ala Glu Leu Gly Thr Ile Lys Asp Pro Tyr Arg Leu

405 410 415

Cys Ser Ile Ser Glu Asp Ser Gly Leu Ser Thr Ala Phe Thr Ile Ala
420 425 430

His Glu Leu Gly His Val Phe Asn Met Pro His Asp Asp Asn Asn Lys
435 440 445

Cys Lys Glu Glu Gly Val Lys Ser Pro Gln His Val Met Ala Pro Thr
450 455 460

Leu Asn Phe Tyr Thr Asn Pro Trp Met Trp Ser Lys Cys Ser Arg Lys
465 470 475 480

Tyr Ile Thr Glu Phe Leu Asp Thr Gly Tyr Gly Glu Cys Leu Leu Asn
485 490 495

Glu Pro Glu Ser Arg Pro Tyr Pro Leu Pro Val Gln Leu Pro Gly Ile
500 505 510

Leu Tyr Asn Val Asn Lys Gln Cys Glu Leu Ile Phe Gly Pro Gly Ser
515 520 525

Gln Val Cys Pro Tyr Met Met Gln Cys Arg Arg Leu Trp Ser Asn Asn
530 535 540

Val Asn Gly Val His Lys Gly Cys Arg Thr Gln His Thr Pro Trp Ala
545 550 555 560

Asp Gly Thr Glu Cys Glu Pro Gly Lys His Cys Lys Tyr Gly Phe Cys
565 570 575

Val Pro Lys Glu Met Asp Val Pro Val Thr Asp Gly Ser Trp Gly Ser
580 585 590

Trp Ser Pro Phe Gly Thr Cys Ser Arg Thr Cys Gly Gly Gly Ile Lys
595 600 605

Thr Ala Ile Arg Glu Cys Asn Arg Pro Glu Pro Lys Asn Gly Gly Lys
610 615 620

Tyr Cys Val Gly Arg Arg Met Lys Phe Lys Ser Cys Asn Thr Glu Pro
625 630 635 640

Cys Leu Lys Gln Lys Arg Asp Phe Arg Asp Glu Gln Cys Ala His Phe
645 650 655

Asp Gly Lys His Phe Asn Ile Asn Gly Leu Leu Pro Asn Val Arg Trp
660 665 670

Val Pro Lys Tyr Ser Gly Ile Leu Met Lys Asp Arg Cys Lys Leu Phe
675 680 685

Cys Arg Val Ala Gly Asn Thr Ala Tyr Tyr Gln Leu Arg Asp Arg Val
690 695 700

Ile Asp Gly Thr Pro Cys Gly Gln Asp Thr Asn Asp Ile Cys Val Gln
705 710 715 720

Thr Leu Ala Gly Thr Pro Asn Thr Val His Tyr Gly Tyr Asn Thr Val

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          755                 760                 765

   Val Arg Ile Pro Ala Gly Ala Thr Asn Ile Asp Val Arg Gln His Ser
       770                 775                 780
 5
   Phe Ser Gly Glu Thr Asp Asp Asp Asn Tyr Leu Ala Leu Ser Ser Ser
   785                 790                 795                 800

   Lys Gly Glu Phe Leu Leu Asn Gly Asn Phe Val Val Thr Met Ala Lys
10                 805                 810                 815

   Arg Glu Ile Arg Ile Gly Asn Ala Val Val Glu Tyr Ser Gly Ser Glu
               820                 825                 830

15 Thr Ala Val Glu Arg Ile Asn Ser Thr Asp Arg Ile Glu Gln Glu Leu
           835                 840                 845

   Leu Leu Gln Val Leu Ser Val Gly Lys Leu Tyr Asn Pro Asp Val Arg
       850                 855                 860
20
   Tyr Ser Phe Asn Ile Pro Ile Glu Asp Lys Pro Gln Gln Phe Tyr Trp
   865                 870                 875                 880

   Asn Ser His Gly Pro Trp Gln Ala Cys Ser Lys Pro Cys Gln Gly Glu
25                 885                 890                 895

   Arg Lys Arg Lys Leu Val Cys Thr Arg Glu Ser Asp Gln Leu Thr Val
               900                 905                 910

30 Ser Asp Gln Arg Cys Asp Arg Leu Pro Gln Pro Gly His Ile Thr Glu
           915                 920                 925

   Pro Cys Gly Thr Gly Cys Asp Leu Arg Trp His Val Ala Ser Arg Ser
       930                 935                 940
35
   Glu Cys Ser Ala Gln Cys Gly Leu Gly Tyr Arg Thr Leu Asp Ile Tyr
   945                 950                 955                 960

   Cys Ala Lys Tyr Ser Arg Leu Asp Gly Lys Thr Glu Lys Val Asp Asp
40                 965                 970                 975

   Gly Phe Cys Ser Ser His Pro Lys Pro Ser Asn Arg Glu Lys Cys Ser
               980                 985                 990

45 Gly Glu Cys Asn Thr Gly Gly Trp Arg Tyr Ser Ala Trp Thr Glu Cys
           995                1000                1005

   Met Lys Ser Cys Asp Gly Gly Thr Gln Arg Arg Arg Ala Ile Cys Val
      1010                1015                1020
50
   Asn Thr Arg Asn Asp Val Leu Asp Asp Ser Lys Cys Thr His Gln Glu
   1025                1030                1035                1040

   Lys Val Thr Ile Gln Arg Cys Ser Glu Phe Pro Cys Pro Gln Trp Lys
55                 1045                1050                1055

   Ser Gly Asp Trp Ser Glu Cys Leu Val Thr Cys Gly Lys Gly His Lys

60

   His Glu Met Ala Ser Trp Gln Ala Gly Pro Trp Val Gln Cys Ser Val
```

1105 1110 1115 1120

Thr Cys Gly Gln Gly Tyr Gln Leu Arg Ala Val Lys Cys Ile Ile Gly
1125 1130 1135

Thr Tyr Met Ser Val Val Asp Asp Asn Asp Cys Asn Ala Ala Thr Arg
1140 1145 1150

Pro Thr Asp Thr Gln Asp Cys Glu Leu Pro Ser Cys His Pro Pro Pro
1155 1160 1165

Ala Ala Pro Glu Thr Arg Arg Ser Thr Tyr Ser Ala Pro Arg Thr Gln
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Trp Arg Phe Gly Ser Trp Thr Pro Cys Ser Ala Thr Cys Gly Lys Gly
1185 1190 1195 1200

Thr Arg Met Arg Tyr Val Ser Cys Arg Asp Glu Asn Gly Ser Val Ala
1205 1210 1215

Asp Glu Ser Ala Cys Ala Thr Leu Pro Arg Pro Val Ala Lys Glu Glu
1220 1225 1230

Cys Ser Val Thr Pro Cys Gly Gln Trp Lys Ala Leu Asp Trp Ser Ser
1235 1240 1245

Cys Ser Val Thr Cys Gly Gln Gly Arg Ala Thr Arg Gln Val Met Cys
1250 1255 1260

Val Asn Tyr Ser Asp His Val Ile Asp Arg Ser Glu Cys Asp Gln Asp
1265 1270 1275 1280

Tyr Ile Pro Glu Thr Asp Gln Asp Cys Ser Met Ser Pro Cys Pro Gln
1285 1290 1295

Arg Thr Pro Asp Ser Gly Leu Ala Gln His Pro Phe Gln Asn Glu Asp
1300 1305 1310

Tyr Arg Pro Arg Ser Ala Ser Pro Ser Arg Thr His Val Leu Gly Gly
1315 1320 1325

Asn Gln Trp Arg Thr Gly Pro Trp Gly Ala Cys Ser Ser Thr Cys Ala
1330 1335 1340

Gly Gly Ser Gln Arg Arg Val Val Val Cys Gln Asp Glu Asn Gly Tyr
1345 1350 1355 1360

Thr Ala Asn Asp Cys Val Glu Arg Ile Lys Pro Asp Ser Gln Arg Ala
1365 1370 1375

Cys Glu Ser Gly Pro Cys Pro Gln Trp Ala Tyr Gly Asn Trp Gly Glu
1380 1385 1390

Cys Thr Lys Leu Cys Gly Gly Gly Ile Arg Thr Arg Leu Val Val Cys
1395 1400 1405

Gln Arg Ser Asn Gly Glu Arg Phe Pro Asp Leu Ser Cys Glu Ile Leu

Gly Arg Gly His Lys Gln Arg Asn Val Tyr Cys Met Ala Lys Asp Gly

```
            1460                1465                1470

   Ser His Leu Glu Ser Asp Tyr Cys Lys His Leu Ala Lys Pro His Gly
          1475                1480                1485

 5
   His Arg Lys Cys Arg Gly Gly Arg Cys Pro Lys Trp Lys Ala Gly Ala
      1490                1495                1500

   Trp Ser Gln Cys Ser Val Ser Cys Gly Arg Gly Val Gln Gln Arg His
10 1505                1510                1515                1520

   Val Gly Cys Gln Ile Gly Thr His Lys Ile Ala Arg Asp Thr Glu Cys
                  1525                1530                1535

15 Asn Pro Tyr Thr Arg Pro Glu Ser Glu Cys Glu Cys Gln Gly Pro Arg
              1540                1545                1550

   Cys Pro Leu Tyr Thr Trp Arg Ala Glu Glu Ser Gln Glu Cys Thr Lys
          1555                1560                1565

20
   Thr Cys Gly Glu Gly Ser Arg Tyr Arg Lys Val Val Cys Val Asp Asp
      1570                1575                1580

   Asn Lys Asn Glu Val His Gly Ala Arg Cys Asp Val Ser Lys Arg Pro
25 1585                1590                1595                1600

   Val Asp Arg Glu Ser Cys Ser Leu Gln Pro Cys Glu Tyr Val Trp Ile
                  1605                1610                1615

30 Thr Gly Glu Trp Ser Glu Cys Ser Val Thr Cys Gly Lys Gly Tyr Lys
              1620                1625                1630

   Gln Arg Leu Val Ser Cys Ser Glu Ile Tyr Thr Gly Lys Glu Asn Tyr
          1635                1640                1645

35
   Glu Tyr Ser Tyr Gln Thr Thr Ile Asn Cys Pro Gly Thr Gln Pro Pro
      1650                1655                1660

   Ser Val His Pro Cys Tyr Leu Arg Glu Cys Pro Val Ser Ala Thr Trp
40 1665                1670                1675                1680

   Arg Val Gly Asn Trp Gly Ser Cys Ser Val Ser Cys Gly Val Gly Val
                  1685                1690                1695

45 Met Gln Arg Ser Val Gln Cys Leu Thr Asn Glu Asp Gln Pro Ser His
              1700                1705                1710

   Leu Cys His Thr Asp Leu Lys Pro Glu Glu Arg Lys Thr Cys Arg Asn
          1715                1720                1725

50
   Val Tyr Asn Cys Glu Leu Pro Gln Asn Cys Lys Glu Val Lys Arg Leu
      1730                1735                1740

   Lys Gly Ala Ser Glu Asp Gly Glu Tyr Phe Leu Met Ile Arg Gly Lys
55 1745                1750                1755                1760

   Leu Leu Lys Ile Phe Cys Ala Gly Met His Ser Asp His Pro Lys Glu
                  1765                1770                1775

60

   Arg Asp Asp Cys Leu Lys Arg Lys Asp Thr Thr Ala Ala Gly Phe Met
```

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      1810                1815                1820

   Ser Phe Gln Lys Ile Arg Ile Asp Leu Thr Ser Met Gln Ile Ile Thr
   1825                1830                1835                1840
 5
   Thr Asp Leu Gln Phe Ala Arg Thr Ser Glu Gly His Pro Val Pro Phe
                  1845                1850                1855

   Ala Thr Ala Gly Asp Cys Tyr Ser Ala Ala Lys Cys Pro Gln Gly Arg
10            1860                1865                1870

   Phe Ser Ile Asn Leu Tyr Gly Thr Gly Leu Ser Leu Thr Glu Ser Ala
         1875                1880                1885

15 Arg Trp Ile Ser Gln Gly Asn Tyr Ala Val Ser Asp Ile Lys Lys Ser
      1890                1895                1900

   Pro Asp Gly Thr Arg Val Val Gly Lys Cys Gly Gly Tyr Cys Gly Lys
   1905                1910                1915                1920
20
   Cys Thr Pro Ser Ser Gly Thr Gly Leu Glu Val Arg Val Leu
                  1925                1930


25
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